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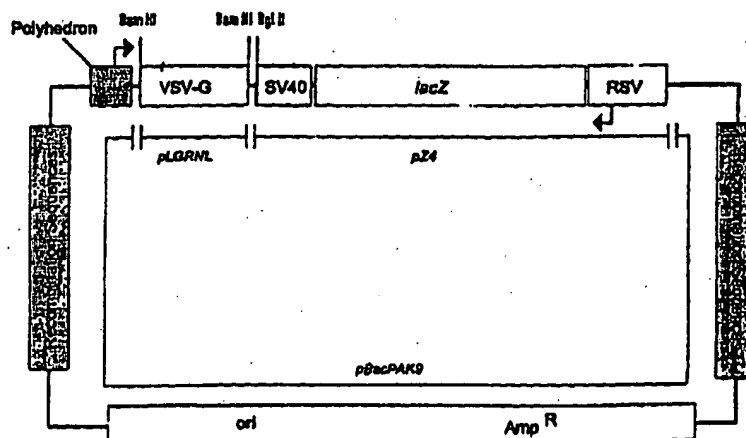


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<p>(21) International Application Number: PCT/US97/16041 (22) International Filing Date: 11 September 1997 (11.09.97) (30) Priority Data: 60/026,297 11 September 1996 (11.09.96) US (71) Applicants (for all designated States except US): THE GENERAL HOSPITAL CORPORATION [US/US]; 55 Fruit Street, Boston, MA 02114 (US). BIOGEN [US/US]; Fourteen Cambridge Center, Cambridge, MA 02142 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): BOYCE, Frederick, M. [-/US]; 103 Beech Street #3, Belmont, MA 02178 (US). BARSOUM, James, G. [-/US]; 6 Moreland Avenue, Lexington, MA 02173 (US). (74) Agents: ELLISON, Eldora, L. et al.; Suite 500, 601 Thirteenth Street, N.W., Washington, DC 20005 (US).</p>		<p>(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published Without international search report and to be republished upon receipt of that report.</p>

(54) Title: NON-MAMMALIAN DNA VIRUS HAVING AN ALTERED COAT PROTEIN

AcMNPV Transfer Plasmid VGZ3



(57) Abstract

Disclosed are methods, nucleic acids, and cells for expressing an exogenous gene in a mammalian cell, involving introducing into the cell a non-mammalian DNA virus (e.g., a baculovirus) having an altered coat protein, the genome of which virus carries an exogenous gene, and growing the cell under conditions such that the gene is expressed. Also disclosed are methods for treating gene deficiency disorders, neurological disorders, or cancers in a mammal by (1) providing to a cell a therapeutically effective amount of a non-mammalian DNA virus having an altered coat protein, the genome of which virus carries an exogenous, therapeutic gene and (2) growing the cell under conditions such that the exogenous gene is expressed in the mammal.

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NON-MAMMALIAN DNA VIRUS HAVING AN ALTERED COAT PROTEIN

Cross-Reference to Related Applications

5 This application claims priority under 35 U.S.C. §119 from U.S. Serial No. 60/026,297, filed September 11, 1996.

Background of the Invention

 This invention relates to the use of a non-mammalian DNA virus to express an exogenous gene in a mammalian cell.

10 Current methods for expressing an exogenous gene in a mammalian cell include the use of mammalian viral vectors, such as those that are derived from retroviruses, adenoviruses, herpes viruses, vaccinia viruses, polio viruses, or adeno-associated viruses. Other methods of expressing an exogenous gene in a mammalian cell include direct injection of DNA, the use of ligand-DNA
15 conjugates, the use of adenovirus-ligand-DNA conjugates, calcium phosphate precipitation, and methods that utilize a liposome- or polycation-DNA complex. In some cases, the liposome- or polycation-DNA complex is able to target the exogenous gene to a specific type of tissue, such as liver tissue.

 Typically, viruses that are used to express desired genes are constructed
20 by removing unwanted characteristics from a virus that is known to infect, and replicate in, a mammalian cell. For example, the genes encoding viral structural proteins and proteins involved in viral replication often are removed to create a defective virus, and a therapeutic gene is then added. This principle has been used to create gene therapy vectors from many types of animal viruses
25 such as retroviruses, adenoviruses, and herpes viruses. This method has also been applied to Sindbis virus, an RNA virus that normally infects mosquitoes but which can replicate in humans, causing a rash and an arthritis syndrome.

 Non-mammalian viruses have been used to express exogenous genes in non-mammalian cells. For example, viruses of the family Baculoviridae
30 (commonly referred to as baculoviruses) have been used to express exogenous genes in insect cells. One of the most studied baculoviruses is *Autographa californica* multiple nuclear polyhedrosis virus (AcMNPV). Although some

genes in insect cells. One of the most studied baculoviruses is *Autographa californica* multiple nuclear polyhedrosis virus (AcMNPV). Although some species of baculoviruses that infect crustacea have been described (Blissard, et al., 1990, Ann. Rev. Entomology 35:127), the normal host range of the

5 baculovirus AcMNPV is limited to the order lepidoptera. Baculoviruses have been reported to enter mammalian cells (Volkman and Goldsmith, 1983, Appl. and Environ. Microbiol. 45:1085-1093; Carbonell and Miller, 1987, Appl. and Environ. Microbiol. 53:1412-1417; Brusca et al., 1986, Intervirology 26:207-222; and Tjia et al., 1983, Virology 125:107-117). Although an early report of

10 baculovirus-mediated gene expression in mammalian cells appeared, the authors later attributed the apparent reporter gene activity to the reporter gene product being carried into the cell after a prolonged incubation of the cell with the virus (Carbonell et al., 1985, J. Virol. 56:153-160; and Carbonell and Miller, 1987, Appl. and Environ. Microbiol. 53:1412-1417). These authors reported that,

15 when the exogenous gene gains access to the cell as part of the baculovirus genome, the exogenous gene is not expressed *de novo*. Subsequent studies have demonstrated baculovirus-mediated gene expression in mammalian cells (Boyce and Bucher, 1996, Proc. Natl. Acad. Sci. 93:2348-2352). In addition to the Baculoviridae, other families of viruses naturally multiply only in invertebrates;

20 some of these viruses are listed in Table 1.

Gene therapy methods are currently being investigated for their usefulness in treating a variety of disorders. Most gene therapy methods involve supplying an exogenous gene to overcome a deficiency in the expression of a gene in the patient. Other gene therapy methods are designed

25 to counteract the effects of a disease. Still other gene therapy methods involve supplying an antisense nucleic acid (e.g., RNA) to inhibit expression of a gene of the host cell (e.g., an oncogene) or expression of a gene from a pathogen (e.g., a virus).

Certain gene therapy methods are being examined for their ability to

30 correct inborn errors of the urea cycle, for example (see, e.g., Wilson et al., 1992, J. Biol. Chem. 267: 11483-11489). The urea cycle is the predominant metabolic pathway by which nitrogen wastes are eliminated from the body.

The steps of the urea cycle are primarily limited to the liver, with the first two steps occurring within hepatic mitochondria. In the first step, carbamoyl phosphate is synthesized in a reaction that is catalyzed by carbamoyl phosphate synthetase I (CPS-I). In the second step, citrulline is formed in a reaction
5 catalyzed by ornithine transcarbamylase (OTC). Citrulline then is transported to the cytoplasm and condensed with aspartate into arginosuccinate by arginosuccinate synthetase (AS). In the next step, arginosuccinate lyase (ASL) cleaves arginosuccinate to produce arginine and fumarate. In the last step of the cycle, arginase converts arginine into ornithine and urea.

10 A deficiency in any of the five enzymes involved in the urea cycle has significant pathological effects, such as lethargy, poor feeding, mental retardation, coma, or death within the neonatal period (see, e.g., Emery et al., 1990, *In: Principles and Practice of Medical Genetics*, Churchill Livingstone, New York). OTC deficiency usually manifests as a lethal hyperammonemic
15 coma within the neonatal period. A deficiency in AS results in citrullinemia which is characterized by high levels of citrulline in the blood. The absence of ASL results in arginosuccinic aciduria (ASA), which results in a variety of conditions including severe neonatal hyperammonemia and mild mental retardation. An absence of arginase results in hyperarginemia which can
20 manifest as progressive spasticity and mental retardation during early childhood. Other currently used therapies for hepatic disorders include dietary restrictions; liver transplantation; and administration of arginine freebase, sodium benzoate, and/or sodium phenylacetate.

Summary of the Invention

25 It has been discovered that a non-mammalian DNA virus carrying an exogenous gene expression construct and having an "altered" coat protein can be used to increase the efficiency with which the non-mammalian DNA virus expresses an exogenous gene in the mammalian cell. For example, expression of vesicular stomatitis virus glycoprotein G (VSV-G) as an altered coat protein
30 on the surface of a virus particle of a baculovirus enhances the ability of the baculovirus to express an exogenous gene (e.g., a therapeutic gene) in a

mammalian cell. While not being obligated to know or disclose the manner in which the invention works, nor being bound by any theory, it is postulated that the enhancement of gene expression is due to an increased ability of a virus particle to gain entry into the cytosol of the mammalian cell.

5 Accordingly, in one aspect, the invention features a method of expressing an exogenous gene in a mammalian cell(s), involving (i) introducing into the cell a non-mammalian DNA virus having an altered coat protein, the genome of which virus carries the exogenous gene under the control of a promoter that induces expression of the exogenous gene in the cell, and (ii) maintaining the
10 cell under conditions such that the exogenous gene is expressed.

 In a second aspect, the invention features a method of treating a gene deficiency disorder in a mammal (e.g., a human or a mouse), involving introducing into a cell (*in vivo* or *ex vivo*) a therapeutically effective amount of a non-mammalian DNA virus having an altered coat protein, the genome of
15 which virus carries an exogenous gene, and maintaining the cell under conditions such that the exogenous gene is expressed in the mammal.

 The invention further features a method for treating a tumor in a mammal, involving introducing into a cancerous cell of the mammal (e.g., a cancerous hepatocyte) a non-mammalian DNA virus (e.g., a baculovirus) having
20 an altered coat protein, the genome of which virus expresses a cancer-therapeutic gene (encoding, e.g., a tumor necrosis factor, thymidine kinase, diphtheria toxin chimera, or cytosine deaminase). The exogenous gene can be expressed in a variety of cells, e.g., hepatocytes; cells of the central nervous system, including neural cells such as neurons from brain, spinal cord, or
25 peripheral nerve; adrenal medullary cells; glial cells; skin cells; spleen cells; muscle cells; kidney cells; and bladder cells. Thus, the invention can be used to treat various cancerous or non-cancerous tumors, including carcinomas (e.g., hepatocellular carcinoma), sarcomas, gliomas, and neuromas. Included within the invention are methods for treating lung, breast, and prostate cancers. Either
30 *in vivo* or *in vitro* methods can be used to introduce the virus into the cell in this aspect of the invention. Preferably, the exogenous gene is operably linked to a promoter that is active in cancerous cells, but not in other cells, of the

mammal. For example, the α -fetoprotein promoter is active in cells of hepatocellular carcinomas and in fetal tissue but it is otherwise not active in mature tissues. Accordingly, the use of such a promoter is preferred for expressing a cancer-therapeutic gene for treating hepatocellular carcinomas.

5 The invention also features a method for treating a neurological disorder (e.g., Parkinson's Disease, Alzheimer's Disease, or disorders resulting from injuries to the central nervous system) in a mammal. The method involves (a) introducing into a cell a therapeutically effective amount of a non-mammalian DNA virus (e.g., a baculovirus) having an altered coat protein, the genome of
10 which virus includes an exogenous gene encoding a therapeutic protein, and (b) maintaining the cell under conditions such that the exogenous gene is expressed in the mammal. Particularly useful exogenous genes include those that encode therapeutic proteins such as nerve growth factor, hypoxanthine guanine phosphoribosyl transferase (HGPRT), tyrosine hydroxylase, dopadecarboxylase,
15 brain-derived neurotrophic factor, basic fibroblast growth factor, sonic hedgehog protein, glial derived neurotrophic factor (GDNF) and RETLI (also known as GDNFR α , GFR-1, and TRN1). Both neuronal and non-neuronal cells (e.g., fibroblasts, myoblasts, and kidney cells) are useful in this aspect of the invention. Such cells can be autologous or heterologous to the treated
20 mammal. Preferably, the cell is autologous to the mammal, as such cells obviate concerns about graft rejection. Preferably, the cell is a primary cell, such as a primary neuronal cell or a primary myoblast.

 In each aspect of the invention, the non-mammalian DNA virus is preferably an "invertebrate virus" (i.e., a virus that infects, and replicates in, an
25 invertebrate). For example, the DNA viruses listed in Table 1 can be engineered to have an altered coat protein(s) and used in the invention. Preferably, the invertebrate DNA virus is a baculovirus, e.g., a nuclear polyhedrosis virus, such as an *Autographa californica* multiple nuclear polyhedrosis virus. If desired, the nuclear polyhedrosis virus may be
30 engineered such that it lacks a functional polyhedrin gene. Either or both the occluded form and budded form of virus (e.g., baculovirus) can be used. Another preferred virus is *Bombyx mori* Nuclear Polyhedrosis Virus (BmNPV).

TABLE 1. NON-MAMMALIAN DNA VIRUSES THAT CAN BE USED IN THE INVENTION.¹

- 5 **I. FAMILY: BACULOVIRUSES BACULOVIRIDAE**
- SUBFAMILY:**
- OCCLUDED BACULOVIRUSES EUBACULOVIRINAE
- Genus:**
- 10 Nuclear polyhedrosis virus (NPV)
- Subgenus:**
- Multiple Nucleocapsid Viruses (MNPV)
- Preferred Species:**
- Autographa californica nuclear polyhedrosis virus (AcMNPV)
- Other Members:**
- 15 Choristoneura fumiferana MNPV (CfMNPV)
- Mamestra brassicae MNPV (MbMNPV)
- Orgyia pseudotsugata MNPV (OpMNPV)
- and approximately 400-500 species
- isolated from seven insect orders and Crustacea.
- 20 **Subgenus:**
- Single Nucleocapsid Viruses (SNPV)
- Preferred Species:**
- Bombyx mori S Nuclear Polyhedrosis Virus (BmNPV)
- Other Members:**
- 25 Heliothis zea SNPV (HzSnpv)
- Trichoplusia ni SNPV (TnSnpv)
- and similar viruses isolated from seven insect
- orders and Crustacea.
- 30 **Genus:**
- Granulosis virus (GV)

¹ These viruses are listed in: "Fifth Report of the International Committee on Taxonomy of Viruses" (ICTV) by Comelia Buchen-Osmond, 1991, Research School of Biological Sciences, Canberra, Australia. Most viruses listed here are available from the American Type Culture Collection.

Preferred Species:

Plodia interpunctella granulosis virus (PiGV)

Other Members:

- 5 Trichoplusia ni granulosis virus (TnGV)
 Pieris brassicae granulosis virus (PbGV)
 Artogeia rapae granulosis virus (ArGV)
 Cydia pomonella granulosis virus (CpGV)
 and similar viruses from about 50 species in the Lepidoptera

SUBFAMILY:

- 10 NON-OCCLUDED BACULOVIRUSES NUDIBACULOVIRINAE

Genus:

Non-occluded baculoviruses (NOB)

Preferred Species:

Heliothis zea NOB (HzNOB)

- 15 Other Members:

Oryctes rhinoceros virus

Additional viruses have been observed in a fungus
 (Strongwellsea magna), a spider, the European crab
 (Carcinus maenas), and the blue crab (Callinectes
 20 sapidus).

II. FAMILY: ICOSAEDRAL CYTOPLASMIC DEOXYRIBOVIRUSES IRIDOVIRIDAEGenus:

Small iridescent Iridovirus insect virus group

Preferred Species:

- 25 Chilo iridescent virus

Other Members:

- | | | |
|----|----------------------------|----------------------------|
| | Insect iridescent virus 1 | Insect iridescent virus 2 |
| | Insect iridescent virus 6 | Insect iridescent virus 9 |
| | Insect iridescent virus 10 | Insect iridescent virus 16 |
| 30 | Insect iridescent virus 17 | Insect iridescent virus 18 |
| | Insect iridescent virus 19 | Insect iridescent virus 20 |
| | Insect iridescent virus 21 | Insect iridescent virus 22 |
| | Insect iridescent virus 23 | Insect iridescent virus 24 |
| | Insect iridescent virus 25 | Insect iridescent virus 26 |
| 35 | Insect iridescent virus 27 | Insect iridescent virus 28 |
| | Insect iridescent virus 29 | Insect iridescent virus 30 |
| | Insect iridescent virus 31 | Insect iridescent virus 32 |

Genus:

Large iridescent Chloriridovirus insect virus group

Preferred Species:

- 5 Mosquito iridescent virus (iridescent virus - type
3, regular strain)

Other Members:

- 10 Insect iridescent virus 3 Insect iridescent virus 4
Insect iridescent virus 5 Insect iridescent virus 7
Insect iridescent virus 8 Insect iridescent virus 11
Insect iridescent virus 12 Insect iridescent virus 13
Insect iridescent virus 14 Insect iridescent virus 15

Putative member:

Chironomus plumosus iridescent

Genus:

- 15 Frog virus group Ranavirus

Preferred Species:

Frog virus 3 (FV3)

Other Members:

- 20 Frog virus 1 Frog virus 2 Frog virus 5
Frog virus 6 Frog virus 7 Frog virus 8
Frog virus 9 Frog virus 10 Frog virus 11
Frog virus 12 Frog virus 13 Frog virus 14
Frog virus 15 Frog virus 16 Frog virus 17
25 Frog virus 18 Frog virus 19 Frog virus 20
Frog virus 21 Frog virus 22 Frog virus 23
Frog virus 24 L2 L4 L5
LT 1 LT 2 LT 3 LT 4
T 21 T 6 T 7 T 8
30 T 9 T 10 T 11 T 12
T 13 T 14 T 15 T 16
T 17 T 18 T 19 T 20
Tadpole edema virus from newts
Tadpole edema virus from Rana catesbriana
35 Tadpole edema virus from Xenopus

Genus:

Lymphocystis disease virus group
Lymphocystisvirus

Preferred Species:

Flounder isolate (LCDV-1)

Other Members:

Lymphocystis disease virus dab isolate (LCDV-2)

5

Putative member:

Octopus vulgaris disease virus

Genus

Goldfish virus group

10

Preferred Species:

Goldfish virus 1 (GFV-1)

Other Member:

Goldfish virus 2 (GF-2)

III. FAMILY: PARVOVIRIDAE**Genus**

15

Insect parvovirus group Densovirus

Preferred Species:

Galleria densovirus

Other Members:

20

Junonia Densovirus
 Agraulis Densovirus
 Bombyx Densovirus
 Aedes Densovirus

Putative Members:

25

Acheta Densovirus	Simulium Densovirus
Diatraea Densovirus	Euxoa Densovirus
Leucorrhinia Densovirus	Periplanata Densovirus
Pieris Densovirus	Sibine Densovirus
PC 84 (parvo-like virus from the crab	
Carcinus mediterraneus)	
Hepatopancreatic parvo-like virus of	
penaeid shrimp	

30

IV. FAMILY: POXVIRUS GROUP POXVIRIDAE**SUBFAMILY:****POXVIRUSES OF VERTEBRATES CHORDOPOXVIRINAE****Genus:****5 Molluscum contagiosum subgroup Molluscipoxvirus****Preferred Species:****Molluscum contagiosum virus****SUBFAMILY:****POXVIRUS OF INSECTS ENTOMOPOXVIRINAE****10 Putative Genus:****Entomopoxvirus A Poxvirus of Coleoptera****Preferred Species:****Poxvirus of Melolontha****Other Members:****15 Coleoptera:****Anomala cuprea****Aphodius tasmaniae****Demodema boranensis****Dermolepida albohirtum****20 Figulus sublaevis****Geotrupes sylvaticus****Putative Genus:****Entomopoxvirus B Poxvirus of Lepidoptera and****25 Orthoptera****Preferred Species:****Poxvirus of Amsacta moorei (Lepidoptera)****Other Members:****Lepidoptera:****30 Acrobasis zelleri****Choristoneura biennis****Choristoneura conflictata****Choristoneura diversum****Chorizagrotis auxiliaris****35 Operophtera brumata****Orthoptera:****Arphia conspersa**

Locusta migratoria
 Melanoplus sanguinipes
 Oedaleus senegalensis
 Schistocerca gregaria

- 5 **Putative Genus:**
 Entomopoxvirus C Poxvirus of Diptera

Preferred Species:
 Poxvirus of Chironomus luridus (Diptera)

- Other Members:
 10 Diptera:
 Aedes aegypti
 Camptochironomus tentans
 Chironomus attenuatus
 Chironomus plumosus
 15 Goeldichironomus holoprasimus

- Other members of family Poxviridae
 Albatrosspox (Avipoxvirus)
 Cotia
 Embu
 20 Marmosetpox
 Marsupialpox (Australian 'quokkas')
 Mule deer poxvirus (Odocoileus hemionus;
 Capripoxvirus)
 Volepox (Microtus oeconomus, Microtus
 25 pennsylvanicus)
 Skunk poxvirus (Mephitis; Orthopoxvirus)

V. GROUP CAULIFLOWER CAULIMOVIRUS MOSAIC VIRUS

- Preferred Member:
 30 Cauliflower mosaic virus (CaMV) (cabbage b, davis
 isolate)

- Other Members:
 35 Blueberry red ringspot (327) Carnation etched ring (182)
 Dahlia mosaic (51) Figwort mosaic
 Horseradish latent Mirabilis mosaic
 Peanut chlorotic streak Soybean chlorotic mottle (331)
 Strawberry vein banding (219) Thistle mottle

- Putative Members:
 Aquilegia necrotic mosaic Cassava vein mosaic
 Cestrum virus Petunia vein clearing

Plantago virus 4

Sonchus mottle

VI. GROUP GEMINIVIRUS**Subgroup I (i.e., Genus)**

Maize streak virus

- 5 Preferred Member:
 Maize streak virus (MSV) (133)

- Other Members:
 Chloris striate mosaic (221)
 Digitaria streak
 10 Miscanthus streak
 Wheat dwarf

- Putative Members:
 Bajra streak
 Bromus striate mosaic
 15 Digitaria striate mosaic
 Oat chlorotic stripe
 Paspalum striate mosaic

Subgroup II (i.e., Genus):

Beet curly top virus

- 20 Preferred Member:
 Beet curly top virus (BCTV)(210)

- Other Members:
 Tomato pseudo-curly top virus
 Bean summer death virus
 25 Tobacco yellow dwarf virus
 Tomato leafroll virus

Subgroup III (i.e., Genus):

Bean golden mosaic virus

- 30 Preferred Member:
 Bean golden mosaic virus (BGMV) (192)

Other Members:

Abutilon mosaic virus
 leaf crumple virus
 yellow mosaic virus

African cassava mosaic virusCotton
 Euphorbia mosaic virusHorsegram
 Indian cassava mosaic virusJatropha

- | | | |
|----|--------------------------------------|-------------------------------|
| | mosaic virus | Limabean golden mosaic virus |
| | Malvaceous chlorosis virus | Melon leaf curl virus |
| | Mungbean yellow mosaic virus | Potato yellow mosaic virus |
| | Rhynchosia mosaic virus | Squash leaf curl virus |
| 5 | Tigre disease virus | Tobacco leaf curl virus |
| | Tomato golden mosaic virus | Tomato leaf curl virus |
| | Tomato yellow dwarf virus | Tomato yellow leaf curl virus |
| | Tomato yellow mosaic virus | Watermelon curly mottle virus |
| | Watermelon chlorotic stunt virus | |
| 10 | Honeysuckle yellow vein mosaic virus | |

Putative Members:

- | | | |
|----|--------------------------------|------------------------------|
| | Cotton leaf curl virus | Cowpea golden mosaic virus |
| | Eggplant yellow mosaic virus | Eupatorium yellow vein virus |
| | Lupin leaf curl virus | Soyabean crinkle leaf virus |
| 15 | Solanum apical leaf curl virus | Wissadula mosaic virus |

VII. FAMILY: dsDNA ALGAL VIRUSES PHYCODNAVIRIDAE**Genus:**

dsdna Phycovirus Phycodnavirus group

Preferred Species:

- 20 Paramecium bursaria chlorella virus - 1 (PBCV - 1)
 Viruses of:
 Paramecium bursaria Chlorella NC64A viruses (NC64A viruses)
 Paramecium bursaria Chlorella Pbi viruses (Pbi viruses)
 Hydra viridis Chlorella viruses (HVCV)

25 Other Members:

Chlorella NC64A viruses (thirty-seven NC64A viruses, including PBCV-1)
 Chlorella virus NE-8D (CV-NE8D; synonym NE-8D)

- | | | | |
|----|---------|------------|---------|
| | CV-NYb1 | CV-CA4B | CV-AL1A |
| | CV-NY2C | CV-NC1D | CV-NC1C |
| 30 | CV-CA1A | CV-CA2A | CV-IL2A |
| | CV-IL2B | CV-IL3A | CV-IL3D |
| | CV-SC1A | CV-SC1B | CV-NC1A |
| | CV-NE8A | CV-AL2C | CV-MA1E |
| | CV-NY2F | CV-CA1D | CV-NC1B |
| 35 | CV-NYs1 | CV-IL5-2s1 | CV-AL2A |
| | CV-MA1D | CV-NY2B | CV-CA4A |
| | CV-NY2A | CV-XZ3A | CV-SH6A |
| | CV-BJ2C | CV-XZ6E | CV-XZ4C |
| | CV-XZ5C | CV-XZ4A | |

- 40 Chlorella Pbi viruses

CVA-1
CVM-1

CVB-1
CVR-1

CVG-1

Hydra viridis Chlorella viruses

5 HVCV-1
HVCV-2
HVCV-3

VIII. FAMILY: POLYDNAVIRUS GROUP POLYDNAVIRIDAE

Genus:
Ichnovirus

10 Preferred Species:
Campoplex sonorensis virus (CsV)

Other Member:
Viruses of Glypta sp.

15 **Genus:**
Bracovirus

Preferred Species:
Cotesia melanoscela virus (CmV)

20 The genome of the non-mammalian DNA virus can be engineered to include one or more genetic elements. In general, these elements are selected based on their ability to facilitate expression of (i) an altered coat protein on the surface of a virus particle, and/or (ii) an exogenous gene in a mammalian cell.

Any transmembrane protein that binds to a target mammalian cell, or that mediates membrane fusion to allow escape from endosomes, can be used as the altered coat protein on the non-mammalian DNA virus. Preferably, the altered coat protein is the polypeptide (preferably a glycosylated version) of a glycoprotein that naturally mediates viral infection of a mammalian cell (e.g., a coat protein of a mammalian virus, such as a lentivirus, and influenza virus, a hepatitis virus, or a rhabdovirus). Other useful altered coat proteins include proteins that bind to a receptor on a mammalian cell and stimulate endocytosis. Examples of suitable altered coat proteins include, but are not limited to, the coat proteins listed in Table 2, which are derived from viruses such as HIV,

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influenza viruses, rhabdoviruses, and human respiratory viruses. An exemplary vesicular stomatitis virus glycoprotein G (VSV-G) is encoded by the plasmid BV-CZPG, the nucleotide sequence of which is shown in Fig. 23. If desired, more than one coat protein can be used as altered coat proteins. For example, a first altered coat protein may be a transmembrane protein that binds to a mammalian cell, and a second coat protein may mediate membrane fusion and escape from endosomes.

TABLE 2. EXAMPLES OF SUITABLE ALTERED COAT PROTEINS

	Viral Coat Protein	Reference
10	Vesicular Stomatitis Virus glycoprotein G	GenBank Accession # M21416 ^a
	Herpes Simplex Virus 1 (KOS) glycoprotein B	GenBank Accession # K01760
	Human Immunodeficiency Virus type 1 gp120	GenBank Accession # U47783
15	Influenza A Virus hemagglutinin	GenBank Accession # U38242
	Human Respiratory Syncytial Virus membrane glycoprotein	GenBank Accession # M86651
	Human Respiratory Syncytial Virus fusion protein	GenBank Accession # D00334
20	Tick-Borne Encephalitis Virus glycoprotein E	GenBank Accession # S72426
	Pseudorabies Virus glycoprotein gH	GenBank Accession # M61196
	Rabies Virus G5803FX glycoprotein	GenBank Accession # U11753
25	Human Rhinovirus 1B viral coat proteins VP1, VP2, and VP3	GenBank Accession # D00239
	Semliki Forest Virus coat proteins E1, E2, and E3	GenBank Accession # Z48163
	Human Immunodeficiency Virus-1 envelope spike protein	Mebatsion et al., 1996, PNAS 93:11366-11370

Herpes Simplex Virus-1 Entry Mediator	Montgomery et al., 1996, Cell 87:427-436
Pseudorabies Virus Glycoprotein gE	Enquist et al., 1994, J. Virol. 68:5275-5279
Herpes Simplex Virus Glycoprotein gB	Norais et al., 1996, J. Virol. 70:7379-7387
Bovine Syncytial Virus Envelope Protein	Renshaw et al., 1991, Gene 105:179-184
Human Foamy Virus (HFV)	GenBank Accession # Y07725
Rabies Virus glycoprotein G	Gaudin et al., 1996, J. Virol. 70:7371-7378

The GenBank accession numbers refer to nucleic acid sequences encoding the viral coat proteins.

In a preferred embodiment, the altered coat protein is produced as a fusion (i.e., chimeric) protein. A particularly useful fusion protein includes (i) a transmembrane polypeptide (e.g., antibodies such as IgM, IgG, and single chain antibodies) fused to (ii) a polypeptide that binds to a mammalian cell (e.g., VCAM, NCAM, integrins, and selectins) or to a growth factor. Included among the suitable transmembrane polypeptides are various coat proteins that naturally exist on the surface of a non-mammalian or mammalian virus particle (e.g., baculovirus gp64, influenza hemagglutinin protein, and Vesicular stomatitis virus glycoprotein G). All or a portion of the transmembrane polypeptide can be used, provided that the polypeptide spans the membrane of the virus particle, such that the polypeptide is anchored in the membrane. Non-viral transmembrane polypeptides also can be used. For example, a membrane-bound receptor can be fused to a polypeptide that binds a mammalian cell and used as the altered coat protein. Preferably, the fusion protein includes a viral coat protein (e.g., gp64) and a targeting molecule (e.g., VSV-G). Fusion polypeptides that include all or a cell-binding portion of a cell adhesion molecule also are included within the invention (e.g., a gp64-VCAM fusion protein).

Typically, the gene encoding the altered coat protein is operably linked to a promoter that is not active in the mammalian cell to be infected with the virus but is active in a non-mammalian cell used to propagate the virus (i.e., a "non-mammalian-active" promoter). By contrast, a mammalian-active promoter is used to drive expression of the exogenous gene of interest (e.g., a therapeutic gene), as is discussed below. Generally, promoters derived from viruses that replicate in non-mammalian cells, but which do not replicate in mammalian cells, are useful as non-mammalian active promoters. For example, when using a baculovirus as the non-mammalian DNA virus, a baculovirus polyhedrin promoter can be used to drive expression of the altered coat protein, since baculoviruses do not replicate in mammalian cells. Other examples of suitable non-mammalian active promoters include p10 promoters, p35 promoters, etl promoters, and gp64 promoters, all of which are active in baculoviruses. When insect cells are used to prepare a virus stock, this non-mammalian-active promoter allows the altered coat protein to be expressed on the surface of the resulting virus particles. Upon infecting a mammalian cell with the non-mammalian DNA virus having an altered coat protein, the polyhedrin promoter is inactive. Examples of suitable non-mammalian-active promoters for driving expression of altered coat proteins include baculoviral polyhedrin promoters (e.g., from pAcAb4 from Pharmingen, Inc.), p10 promoters (e.g., from pAcAb4 from Pharmingen, Inc.), p39 promoters (see Xu et al., 1995, J. Virol. 69:2912-2917), gp64 promoters (including TATA-independent promoters; see Kogan et al., 1995, J. Virol. 69:1452-1461), baculoviral IE1 promoters (see Jarvis et al., 1996, Prot. Expr. Purif. 8:191-203), and *Drosophila* alcohol dehydrogenase promoters (see Heberlein et al., 1995, Cell 41:965-977).

If desired, the non-mammalian-active promoter that is operably linked to the gene encoding the altered coat protein can be an inducible promoter that is activated in the non-mammalian cell in which the virus is propagated. Examples of suitable inducible promoters include promoters based on progesterone receptor mutants (Wang et al., 1994, Proc. Natl. Acad. Sci. 91:8180-8184), tetracycline-inducible promoters (Gossen et al., 1995, Science

268:1766-1760; 1992, Proc. Natl. Acad. Sci. 89:5547-5551, available from Clontech, Inc.), rapamycin-inducible promoters (Rivera et al., 1996, Nat. Med. 2:1028-1032), and ecdysone-inducible promoters (No et al., 1996, Proc. Natl. Acad. Sci. 93:3346-3351).

5 In principle, an inducible promoter that can be activated in either a non-mammalian or mammalian cell can be used in this embodiment of the invention, although in practice an inducer of the promoter typically would be added to the non-mammalian cell in which the virus is propagated, rather than the mammalian cell in which the exogenous gene is expressed. As an example,
10 a gene encoding an altered coat protein can be operably linked to a promoter that is inducible by ecdysone (No et al., 1996, Proc. Natl. Acad. Sci. 93:3346-3351). In this case, the genome of the non-mammalian DNA virus is engineered to include a paired ecdysone response element operably linked to the gene encoding the altered coat protein. Expression of a heterodimeric ecdysone
15 receptor in the presence of ecdysone (or an ecdysone analog) that is added to the cell activates gene expression from a promoter that is operably linked to a gene encoding an altered coat protein. The use of an inducible promoter to drive expression of the gene encoding the altered coat protein offers the advantage of providing an additional mechanism for controlling expression of
20 the altered coat protein.

 The genome of the non-mammalian DNA virus can be engineered to include additional genetic elements, such as a mammalian-active promoter of a long-terminal repeat of a transposable element or a retrovirus (e.g., Rous Sarcoma Virus); an inverted terminal repeat of an adeno-associated virus and an
25 adeno-associated rep gene; and/or a cell-immortalizing sequence, such as the SV40 T antigen or c-myc. If desired, the genome of the non-mammalian DNA virus can include an origin of replication that functions in a mammalian cell (e.g., an Epstein Barr Virus (EBV) origin of replication or a mammalian origin of replication). Examples of mammalian origins of replication include
30 sequences near the dihydrofolate reductase gene (Burhans et al., 1990, Cell 62:955-965), the β -globin gene (Kitsberg et al., 1993, Cell 366:588-590), the adenosine deaminase gene (Carroll et al., 1993, Mol. Cell. Biol. 13:2927-2981),

and other human sequences (see Krysan et al., 1989, Mol. Cell. Biol. 9:1026-1033). If desired, the origin of replication can be used in conjunction with a factor that promotes replication of autonomous elements, such as the EBNA1 gene from EBV. The genome of the non-mammalian DNA virus used in the invention can include a polyadenylation signal and an RNA splicing signal that functions in mammalian cells (i.e., a "mammalian RNA splicing signal"), positioned for proper processing of the product of the exogenous gene. In addition, the virus may be engineered to encode a signal sequence for proper targeting of the gene product.

10 The exogenous gene that is to be expressed in a mammalian cell is operably linked to a "mammalian-active" promoter (i.e., a promoter that directs transcription in a mammalian cell). Where cell-type specific expression of the exogenous gene is desired, the exogenous gene in the genome of the virus can be operably linked to a mammalian-active, cell-type-specific promoter, such as
15 a promoter that is specific for liver cells, brain cells (e.g., neuronal cells), glial cells, Schwann cells, lung cells, kidney cells, spleen cells, muscle cells, or skin cells. For example, a liver cell-specific promoter can include a promoter of a gene encoding albumin, α -1-antitrypsin, pyruvate kinase, phosphoenol pyruvate carboxykinase, transferrin, transthyretin, α -fetoprotein, α -fibrinogen, or β -
20 fibrinogen. Alternatively, a hepatitis virus promoter (e.g., hepatitis A, B, C, or D viral promoter) can be used. If desired, a hepatitis B viral enhancer may be used in conjunction with a hepatitis B viral promoter. Preferably, an albumin promoter is used. An α -fetoprotein promoter is particularly useful for driving expression of an exogenous gene when the invention is used to express a gene
25 for treating a hepatocellular carcinoma. Other preferred liver-specific promoters include promoters of the genes encoding the low density lipoprotein receptor, α 2-macroglobulin, α 1-antichymotrypsin, α 2-HS glycoprotein, haptoglobin, ceruloplasmin, plasminogen, complement proteins (C1q, C1r, C2, C3, C4, C5, C6, C8, C9, complement Factor I and Factor H), C3 complement activator, β -
30 lipoprotein, and α 1-acid glycoprotein. For expression of an exogenous gene specifically in neuronal cells, a neuron-specific enolase promoter can be used (see Forss-Petter et al., 1990, Neuron 5: 187-197). For expression of an

exogenous gene in dopaminergic neurons, a tyrosine hydroxylase promoter can be used. For expression in pituitary cells, a pituitary-specific promoter such as POMC may be useful (Hammer et al., 1990, Mol. Endocrinol. 4:1689-97).

Typically, the promoter that is operably linked to the exogenous gene is not
5 identical to the promoter that is operably linked to the gene encoding an altered coat protein.

Promoters that are inducible by external stimuli also can be used for driving expression of the exogenous gene. Such promoters provide a convenient means for controlling expression of the exogenous gene in a cell of
10 a cell culture or within a mammal. Preferred inducible promoters include enkephalin promoters (e.g., the human enkephalin promoter), metallothionein promoters, mouse mammary tumor virus promoters, promoters based on progesterone receptor mutants, tetracycline-inducible promoters, rapamycin-inducible promoters, and ecdysone-inducible promoters. Methods for inducing
15 gene expression from each of these promoters are known in the art.

Essentially any mammalian cell can be used in the invention; preferably, the mammalian cell is a human cell. The cell can be a primary cell (e.g., a primary hepatocyte, primary neuronal cell, or primary myoblast) or it may be a cell of an established cell line. It is not necessary that the cell be
20 capable of undergoing cell division; a terminally differentiated cell can be used in the invention. If desired, the virus can be introduced into a primary cell approximately 24 hours after plating of the primary cell to maximize the efficiency of infection. Preferably, the mammalian cell is a liver-derived cell, such as a HepG2 cell, a Hep3B cell, a Huh-7 cell, an FTO2B cell, a Hepa1-6
25 cell, or an SK-Hep-1 cell) or a Kupffer cell; a kidney cell, such as a cell of the kidney cell line 293, a PC12 cell (e.g., a differentiated PC12 cell induced by nerve growth factor), a COS cell (e.g., a COS7 cell), or a Vero cell (an African green monkey kidney cell); a neuronal cell, such as a fetal neuronal cell, cortical pyramidal cell, mitral cell, a granule cell, or a brain cell (e.g., a cell of
30 the cerebral cortex; an astrocyte; a glial cell; a Schwann cell); a muscle cell, such as a myoblast or myotube (e.g., a C₂C₁₂ cell); an embryonic stem cell, a spleen cell (e.g., a macrophage or lymphocyte); an epithelial cell, such as a

HeLa cell (a human cervical carcinoma epithelial line); a fibroblast, such as an NIH3T3 cell; an endothelial cell; a WISH cell; an A549 cell; or a bone marrow stem cell. Other preferred mammalian cells include CHO/dhfr cells, Ramos, Jurkat, HL60, and K-562 cells.

5 The virus can be introduced into a cell *in vitro* or *in vivo*. Where the virus is introduced into a cell *in vitro*, the infected cell can subsequently be introduced into a mammal, if desired. Accordingly, expression of the exogenous gene can be accomplished by maintaining the cell *in vitro*, *in vivo*, or *in vitro* and *in vivo*, sequentially. Similarly, where the invention is used to
10 express an exogenous gene in more than one cell, a combination of *in vitro* and *in vivo* methods may be used to introduce the gene into more than one mammalian cell.

 If desired, the virus can be introduced into the cell by administering the virus to a mammal that carries the cell. For example, the virus can be
15 administered to a mammal by subcutaneous, intravascular, or intraperitoneal injection. If desired, a slow-release device, such as an implantable pump, may be used to facilitate delivery of the virus to cells of the mammal. A particular cell type within the mammal can be targeted by modulating the amount of the virus administered to the mammal and by controlling the method of delivery.
20 For example, intravascular administration of the virus to the portal, splenic, or mesenteric veins or to the hepatic artery may be used to facilitate targeting the virus to liver cells. In another method, the virus may be administered to cells or an organ of a donor individual (human or non-human) prior to transplantation of the cells or organ to a recipient.

25 In a preferred method of administration, the virus is administered to a tissue or organ containing the targeted cells of the mammal. Such administration can be accomplished by injecting a solution containing the virus into a tissue, such as skin, brain (e.g., the cerebral cortex), kidney, bladder, liver, spleen, muscle, thyroid, thymus, lung, or colon tissue. Alternatively, or
30 in addition, administration can be accomplished by perfusing an organ with a solution containing the virus, according to conventional perfusion protocols.

In another preferred method, the virus is administered intranasally, e.g., by applying a solution of the virus to the nasal mucosa of a mammal. This method of administration can be used to facilitate retrograde transportation of the virus into the brain. This method thus provides a means for delivering the virus to brain cells, (e.g., mitral and granule neuronal cells of the olfactory bulb) without subjecting the mammal to surgery.

In an alternative method for using the virus to express an exogenous gene in the brain, the virus is delivered to the brain by osmotic shock according to conventional methods for inducing osmotic shock.

Where the cell is maintained under *in vitro* conditions, conventional tissue culture conditions and methods may be used. In a preferred method, the cell is maintained on a substrate that contains collagen, such as Type I collagen or rat tail collagen, or a matrix containing laminin. As an alternative to, or in addition to, maintaining the cell under *in vitro* conditions, the cell can be maintained under *in vivo* conditions (e.g., in a human). Implantable versions of collagen substrates are also suitable for maintaining the virus-infected cells under *in vivo* conditions in practicing the invention (see, e.g., Hubbell et al., 1995, Bio/Technology 13:565-576 and Langer and Vacanti, 1993, Science 260: 920-925).

The invention can be used to express a variety of exogenous genes encoding gene products such as a polypeptides or proteins, antisense RNAs, and catalytic RNAs. If desired, the gene product (e.g., protein or RNA) can be purified from the mammalian cell. Thus, the invention can be used in the manufacture of a wide variety of proteins that are useful in the fields of biology and medicine.

Where the invention is used to express an antisense RNA, the preferred antisense RNA is complementary to a nucleic acid (e.g., an mRNA) of a pathogen of the mammalian cell (e.g., a virus, a bacterium, or a fungus). For example, the invention can be used in a method of treating a hepatitis viral infection by expressing an antisense RNA that hybridizes to an mRNA of an essential hepatitis virus gene product (e.g., a polymerase mRNA). Other preferred antisense RNAs include those that are complementary to a naturally-

occurring gene in the cell, which gene is expressed at an undesirably high level. For example, an antisense RNA can be designed to inhibit expression of an oncogene in a mammalian cell. Similarly, the virus can be used to express a catalytic RNA (i.e., a ribozyme) that inhibits expression of a target gene in the
5 cell by hydrolyzing an mRNA encoding the targeted gene product. Antisense RNAs and catalytic RNAs can be designed by employing conventional criteria.

If desired, the invention can be used to express a dominant negative mutant in a mammalian cell. For example, viral assembly in a cell can be inhibited or prevented by expressing in that cell a dominant negative mutant of
10 a viral capsid protein (see, e.g., Scaglioni et al., 1994, Virology 205:112-120; Scaglioni et al., 1996, Hepatology 24:1010-1017; and Scaglioni et al., 1997, J. Virol. 71:345-353).

The invention can be used to express any of various "therapeutic" genes in a cell. A "therapeutic" gene is one that, when expressed, confers a beneficial
15 effect on the cell or tissue in which it is present, or on a mammal in which the gene is expressed. Examples of "beneficial effects" include amelioration of a sign or symptom of a condition or disease, prevention or inhibition of a condition or disease, or conferral of a desirable characteristic. Included among the therapeutic genes are those genes that correct a gene deficiency disorder in
20 a cell or mammal. For example, carbamoyl synthetase I can correct a gene deficiency disorder when it is expressed in a cell that previously failed to express, or expressed insufficient levels of, carbamoyl synthetase I.

"Correction" of a gene deficiency disorder need not be equivalent to curing a patient suffering from a disorder. All that is required is conferral of a
25 beneficial effect, including even temporary amelioration of signs or symptoms of the disorder. Also included are genes that are expressed in one cell, yet which confer a beneficial effect on a second cell. For example, a gene encoding insulin can be expressed in a pancreatic cell, from which the insulin is then secreted to exert an effect on other cells of the mammal. Other
30 therapeutic genes include sequences that encode antisense RNAs nucleic acid that inhibit transcription or translation of a gene that is expressed at an undesirably high level. For example, an antisense gene that inhibits expression

of a gene encoding an oncogenic protein is considered a therapeutic gene.

"Cancer therapeutic" genes are those genes that confer a beneficial effect on a cancerous cell or a mammal suffering from cancer. Particularly useful cancer therapeutic genes include the p53 gene, a herpes simplex virus thymidine kinase gene, and an antisense gene that is complementary to an oncogene.

The invention can be used to express a therapeutic gene in order to treat a gene deficiency disorder. Particularly appropriate genes for expression include those genes that are thought to be expressed at a less than normal level in the target cells of the subject mammal. Particularly useful gene products include carbamoyl synthetase I, ornithine transcarbamylase, arginosuccinate synthetase, arginosuccinate lyase, and arginase. Other desirable gene products include fumarylacetoacetate hydrolase, phenylalanine hydroxylase, alpha-1 antitrypsin, glucose-6-phosphatase, low-density-lipoprotein receptor, porphobilinogen deaminase, factor VIII, factor IX, cystathione β -synthase, branched chain ketoacid decarboxylase, albumin, isovaleryl-CoA dehydrogenase, propionyl CoA carboxylase, methyl malonyl CoA mutase, glutaryl CoA dehydrogenase, insulin, β -glucosidase, pyruvate carboxylase, hepatic phosphorylase, phosphorylase kinase, glycine decarboxylase (also referred to as P-protein), H-protein, T-protein, Menkes disease copper-transporting ATPase, Wilson's disease copper-transporting ATPase, and CFTR (e.g., for treating cystic fibrosis).

The invention can also be used to express in a mammalian cell a gene that is expected to have a biological effect in mammals but not in insects (i.e., a "mammal-specific" gene). For example, a baculovirus genome can be used to express a mammalian myoD gene and thereby produce muscle proteins; such a gene would be expected to have a biological effect in mammalian cells but not insect cells. Other examples of mammal-specific genes include, but are not limited to, transcription factors that function in mammalian, but not insect, cells. For example, the transcription factors c/ebp-alpha and chop10 will activate liver cell differentiation pathways when expressed from an insect genome (e.g., a baculovirus genome) in a mammalian cell. In contrast,

expression of these mammal-specific transcription factors in an insect cell would be expected to have a minimal, or no, effect on the insect cell.

If desired, the vectors of the invention can be used to propagate genetic constructs in non-mammalian (e.g., insect) cells, with the advantage of inhibiting DNA methylation of the product. It has been observed that a promoter may become methylated in cell lines or tissues in which it is not normally expressed, and that such methylation is inhibitory to proper tissue specific expression (Okuse et al., 1997, Brain Res. Mol. Brain Res. 46:197-207; Kudo et al., 1995, J. Biol. Chem. 270:13298-13302). For example, a neural promoter may become methylated in a non-neural mammalian cell. By using, for example, insect cells (e.g., Sf9 cells) to propagate a baculovirus carrying an exogenous gene and a mammalian promoter (e.g., a neural promoter), the invention provides a means for inhibiting DNA methylation of the promoter prior to administration of the baculovirus and exogenous gene to the mammalian cell in which the exogenous gene will be expressed (e.g., a neural cell).

Definitions

By "non-mammalian" DNA virus is meant a virus that has a DNA genome (rather than RNA) and which is naturally incapable of replicating in a vertebrate, and specifically a mammalian, cell. Included are insect viruses (e.g., baculoviruses), amphibian viruses, plant viruses, and fungal viruses. Viruses that naturally replicate in prokaryotes are excluded from this definition. Examples of viruses that are useful in practicing the invention are listed in Table 1. As used herein, a "genome" can include all or some of the nucleic acid sequences present in a naturally-occurring non-mammalian DNA virus. If desired, genes or sequences can be removed from the virus genome or disabled (e.g., by mutagenesis), provided that the virus retains, or is engineered to retain, its ability to express an exogenous gene in a mammalian cell. For example, the virus can be engineered such that it lacks a functional polyhedrin gene. Such a virus can be produced by deleting all or a portion of the polyhedrin gene from a virus genome (e.g., a baculovirus genome) or by introducing mutations (e.g.,

a frameshift mutation) into the polyhedrin gene so that the activity of the gene product is inhibited.

By "insect" DNA virus is meant a virus that has a DNA genome and which is naturally capable of replicating in an insect cell (e.g., Baculoviridae, Iridoviridae, Poxviridae, Polydnaviridae, Densoviridae, Caulimoviridae, and Phycodnaviridae).

By "exogenous" gene or promoter is meant any gene or promoter that is not normally part of the non-mammalian DNA virus (e.g., baculovirus) genome. Such genes include those genes that normally are present in the mammalian cell to be infected; also included are genes that are not normally present in the mammalian cell to be infected (e.g., related and unrelated genes of other cells or species). As used herein, the term "exogenous gene" excludes a gene encoding an "altered coat protein."

By "altered coat protein" is meant any polypeptide that (i) is engineered to be expressed on the surface of a virus particle, (ii) is not naturally present on the surface of the non-mammalian DNA virus used to infect a mammalian cell, and (iii) allows entry to a mammalian cell by binding to the cell and/or facilitating escape from the mammalian endosome into the cytosol of the cell. Typically, a gene encoding an altered coat protein is incorporated into the genome of the non-mammalian DNA virus used in the invention. If desired, a virus genome can be constructed such that the virus expresses a polypeptide that binds a mammalian receptor or counterreceptor on a mammalian cell. An altered coat protein can include all or a portion of a coat protein of a "mammalian" virus, i.e., a virus that naturally infects and replicates in a mammalian cell (e.g., an influenza virus). If desired, the altered coat protein can be a "fusion protein," i.e., an engineered protein that includes part or all of two (or more) distinct proteins derived from one or multiple distinct sources (e.g., proteins of different species). Typically, a fusion protein used in the invention includes (i) a polypeptide that has a transmembrane region of a transmembrane protein (e.g., baculovirus gp64) fused to (ii) a polypeptide that binds a mammalian cell (e.g., an extracellular domain of VSV-G).

Although the term "altered" is used in reference to the coat protein (because it is altered in the sense that it is expressed on the surface of a virus particle on which it is not normally found), the protein itself need not differ in sequence or structure from a wild-type version of the protein. Thus, a wild-type transmembrane protein that binds a mammalian cell can be used as the altered coat protein (e.g., a wild-type influenza virus hemagglutinin protein). Indeed, wild-type proteins are preferred. Nonetheless, non-wild-type proteins also can be used as the "altered" coat protein, provided that the non-wild-type coat protein retains the ability to bind to a mammalian cell. Examples of non-wild-type proteins include truncated proteins, mutant proteins (e.g., deletion mutants), and conservative variations of transmembrane polypeptides that bind a mammalian cell.

"Conservative variation" denotes the replacement of an amino acid residue by another, functionally similar, residue. Examples of conservative variations include the substitution of one hydrophobic residue, such as alanine, isoleucine, valine, leucine, or methionine, for another, or the substitution of one polar residue for another, such as the substitution of arginine for lysine, glutamic acid for aspartic acid, or glutamine for asparagine, and the like. The term "conservative variation" also includes the use of a substituted amino acid (i.e., a modified amino acid, such as Hydroxylysine) in place of an unsubstituted parent amino acid.

By "positioned for expression" is meant that the DNA sequence that includes the reference gene (e.g., the exogenous gene) is positioned adjacent to a DNA sequence that directs transcription of the DNA and, if desired, translation of the RNA (i.e., facilitates the production of the desired gene product).

By "promoter" is meant at least a minimal sequence sufficient to direct transcription. A "mammalian-active" promoter is one that is capable of directing transcription in a mammalian cell. The term "mammalian-active" promoter includes promoters that are derived from the genome of a mammal, i.e., "mammalian promoters," and promoters of viruses that are naturally capable of directing transcription in mammals (e.g., an MMTV promoter).

Other promoters that are useful in the invention include those promoters that are sufficient to render promoter-dependent gene expression controllable for cell-type specificity, cell-stage specificity, or tissue-specificity (e.g., liver-specific promoters), and those promoters that are "inducible" by external signals or agents (e.g., metallothionein, MMTV, and pENK promoters); such elements can be located in the 5' or 3' regions of the native gene. The promoter sequence can be one that does not occur in nature, so long as it functions in a mammalian cell. An "inducible" promoter is a promoter that, (a) in the absence of an inducer, does not direct expression, or directs low levels of expression, of a gene to which the inducible promoter is operably linked; or (b) exhibits a low level of expression in the presence of a regulating factor that, when removed, allows high-level expression from the promoter (e.g., the *tet* system). In the presence of an inducer, an inducible promoter directs transcription at an increased level.

By "operably linked" is meant that a gene and a regulatory sequence(s) (e.g., a promoter) are connected in such a way as to permit gene expression when the appropriate molecules (e.g., transcriptional activator proteins) are bound to the regulatory sequence(s).

By "cell-immortalizing sequence" is meant a nucleic acid that, when present in a mammalian cell, is capable of transforming the cell for prolonged inhibition of senescence. Included are SV40 T-antigen, *c-myc*, telomerase, and E1A.

By "antisense" nucleic acid is meant a nucleic acid molecule (i.e., RNA) that is complementary (i.e., able to hybridize) to all or a portion of a target nucleic acid (e.g., a gene or mRNA) that encodes a polypeptide of interest. If desired, conventional methods can be used to produce an antisense nucleic acid that contains desirable modifications. For example, a phosphorothioate oligonucleotide can be used as the antisense nucleic acid in order to inhibit degradation of the antisense oligonucleotide by nucleases *in vivo*. Where the antisense nucleic acid is complementary to only a portion of the target nucleic acid encoding the polypeptide to be inhibited, the antisense nucleic acid should hybridize close enough to some critical portion of the target

nucleic acid (e.g., in the translation control region of the non-coding sequence, or at the 5' end of the coding sequence) such that it inhibits translation of a functional polypeptide (i.e., a polypeptide that carries out an activity that one wishes to inhibit (e.g., an enzymatic activity)). Typically, this means that the antisense nucleic acid should be complementary to a sequence that is within the 5' half or third of a target mRNA to which the antisense nucleic acid hybridizes. As used herein, an "antisense gene" is a nucleic acid that is transcribed into an antisense RNA. Typically, such an antisense gene includes all or a portion of the target nucleic acid, but the antisense gene is operably linked to a promoter such that the orientation of the antisense gene is opposite to the orientation of the sequence in the naturally-occurring gene.

Use

The invention is useful for expressing an exogenous gene(s) in a mammalian cell *in vitro* or *in vivo* (e.g., a HepG2 cell). This method can be employed in the manufacture of proteins to be purified, such as proteins that are administered as pharmaceutical agents (e.g., insulin). The virus of the invention can also be used therapeutically. For example, the invention can be used to express in a patient a gene encoding a protein that corrects a deficiency in gene expression. In alternative methods of therapy, the invention can be used to express any protein, antisense RNA, or catalytic RNA in a cell.

The non-mammalian viral expression system of the invention offers several advantages. The altered coat protein on the virus enhances the ability of the non-mammalian DNA virus to infect and express a gene in a mammalian cell. Such a coat protein also can be used to confer cell-type specificity on the engineered virus. For example, expression of CD4⁺ on a cell enhances the ability of a virus expressing an HIV envelope gp120 protein to infect such CD4⁺ cells (Mebatsion et al., 1996, Proc. Natl. Acad. Sci. 93:11366-11370).

The invention allows for *de novo* expression of an exogenous gene; thus, detection of the exogenous protein (e.g., β -galactosidase) in an infected cell represents protein that was actually synthesized in the infected cell, as opposed to protein that is carried along with the virus aberrantly. Because the

non-mammalian viruses used in the invention are not normally pathogenic to humans and do not replicate in mammalian cells, concerns about safe handling of these viruses are minimized. Similarly, because the majority of naturally-occurring viral promoters are not normally active in a mammalian cell, production of undesired viral proteins is minimized. While traditional gene therapy vectors are based upon defective viruses that are propagated with helper virus or on a packaging line, the invention employs a virus that is not defective for growth on insect cells for purposes of virus propagation, but is intrinsically, and desirably, defective for growth on mammalian cells. Accordingly, in contrast to some mammalian virus-based gene therapy methods, the non-mammalian virus-based methods of the invention should not provoke a host immune response to proteins expressed by the virus in the mammalian cells.

The non-mammalian virus used in the invention can be propagated with cells grown in serum-free media, eliminating the risk of adventitious infectious agents occasionally present in the serum contaminating a virus preparation. In addition, the use of serum-free media eliminates a significant expense faced by users of mammalian viruses. Certain non-mammalian viruses, such as baculoviruses, can be grown to a high titer (i.e., 10^8 pfu/ml). Generally, the large virus genomes that can be used in the invention (e.g., the baculovirus genome at 130 kbp) can accept large exogenous DNA molecules (e.g., 100 kb). In certain embodiments, the invention employs a virus whose genome has been engineered to contain an exogenous origin of replication (e.g., the EBV *oriP*). The presence of such sequences on the virus genome allows episomal replication of the virus, increasing persistence in the cell. Where the invention is used in the manufacture of proteins to be purified from the cell, the invention offers the advantage that it employs a mammalian expression system. Accordingly, one can expect proper post-translational processing and modification (e.g., glycosylation) of the product of the exogenous gene.

Other features and advantages of the invention will be apparent from the following detailed description, and from the claims.

Brief Description of the Drawings

Fig. 1 is a schematic representation of the AcMNPV RSV-lacZ transfer plasmid pZ4.

Fig. 2 is a schematic representation of the occluded AcMNPV RSV-lacZ transfer plasmid pZ5.

Fig. 3 is a schematic representation of the episomal transfer plasmid pZ-EBV#1, a chimera of baculovirus and Epstein Barr Virus sequences. A virus produced with this transfer plasmid is capable of replicating in a mammalian cell.

Fig. 4A is a schematic representation of a transfer plasmid that allows excision of a gene cassette. Fig. 4B is a schematic representation of the gene cassette excised by the transfer plasmid of Fig. 4A. Excision of the gene cassette is mediated by cre-lox recombination. This strategy allows persistence of an exogenous gene in the absence of viral sequences.

Fig. 5 is a schematic representation of the transfer plasmid, pBV-AVneo, a chimera of baculovirus and Adeno-associated virus sequences. This plasmid is capable of integrating into the genome of the infected cell.

Fig. 6 is a schematic representation of the AcMNPV transfer plasmid pCMV-BV.

Fig. 7 is a schematic representation of the AcMNPV transfer plasmid pCMVZ-BV.

Fig. 8 is a schematic representation of the AcMNPV transfer plasmid pAct-BV.

Fig. 9 is a schematic representation of the AcMNPV transfer plasmid pAZ-BV.

Fig. 10 is a schematic representation of the AcMNPV transfer plasmid pIE45-BV.

Fig. 11 is a schematic representation of the AcMNPV transfer plasmid pNSE4-BV.

Fig. 12 is a schematic representation of the AcMNPV transfer plasmid pTH/SV40/BP9.

Fig. 13 is a schematic representation of the AcMNPV transfer plasmid pTH-Lac/BP9.

Figs. 14A-D are photographs of cells that were stained with X-gal one day post-infection with an AcMNPV virus containing a RSV-*lacZ* cassette.

5 Cells expressing the *lacZ* gene stain darkly with X-gal. Fig. 14A is a photograph of a typical field of HepG2 cells infected at a multiplicity of infection of 15. Fig. 14B is a photograph of a typical field of HepG2 cells infected at a multiplicity of infection of 125; over 25% of the cells were stained. Fig. 14C is a typical field of Sk-Hep-1 cells infected at a multiplicity
10 of infection of 125, showing no positively-stained cells. Fig. 14D is a less typical field of Sk-Hep-1 cells infected at a multiplicity of infection of 125 showing a positively-stained cell. Bar = 55 μ m.

Fig. 15 is a photograph of cells obtained following baculovirus-mediated gene transfer into primary cultures of rat hepatocytes. Over 70% of
15 the cells were stained blue.

Fig. 16 is a graph displaying the dose-dependence of baculovirus-mediated gene transfer. Here, 10^6 HepG2 cells were seeded into 60 mm petri dishes, and one day later the cells were exposed to the indicated dose of an AcMNPV virus containing a RSV-*lacZ* cassette (viral titer = 1.4×10^9 pfu/ml).
20 At one day post-infection, the cells were harvested, and extracts were prepared and assayed for β -galactosidase enzyme activity. Extract activity is expressed in units of β -galactosidase activity as previously defined (Norton and Coffin, 1985, Mol. Cell. Biol. 5:281-290). Enzyme activity was normalized for the protein content of each extract. Each point is the average of three independent
25 assays, with the error bars representing the standard deviation.

Fig. 17 is a graphic representation of results obtained in a time course of baculovirus-mediated expression. HepG2 cells were infected with AcMNPV virus containing a RSV-*lacZ* cassette (multiplicity of infection = 15) at time zero. After one hour, the medium containing the virus was removed and
30 replaced with fresh medium. Infected cells were harvested at the indicated time points and assayed for β -galactosidase activity as is described above. Each plotted point is expressed as the average of three independent assays, with the

error bars representing the standard deviation. Expression from the virus peaked 12-24 hours post-infection and declined thereafter when normalized to total cellular protein.

Fig. 18 is a schematic representation of the AcMNPV transfer plasmid VSVG/BP9.

Fig. 19 is a schematic representation of the AcMNPV transfer plasmid VGZ3.

Fig. 20 is a schematic representation of a budding baculovirus having an altered coat protein. The natural baculovirus cell surface protein (gp64) and the VSV-G protein are represented by "gp64" and "VSV G."

Fig. 21 is a schematic representation of various baculoviral transfer vectors, in which an exogenous gene is operably linked to a viral or mammalian promoter.

Fig. 22 is a graphic representation of the relative transduction efficiencies of Z4 and VGZ3 in HeLa and HepG2 cells. HeLa and HepG2 cells were treated with the VSV G-lacking baculovirus Z4 or the VSV G-containing baculovirus VGZ3 at multiplicities of infection of 1, 10, and 100. Expression of the lacZ gene was determined on the following day by a *in vitro* chemiluminescence assay. --, HepG2 cells treated with VGZ3; -o-, HepG2 treated with Z4; -■-, HeLa treated with VGZ3; -□-, HeLa treated with Z4.

Fig. 23 is a listing of the nucleotide sequence of plasmid BV-CZPG, which encodes a vesicular stomatitis virus G glycoprotein.

Detailed Description

Genetic Manipulation of Viruses

In contrast to conventional gene expression methods, the invention involves modifying non-mammalian DNA viruses that do not naturally infect and replicate in mammalian cells. Thus, the invention is based on the addition of new properties to a non-mammalian DNA virus that allow it to deliver a gene to a mammalian cell and direct gene expression within the mammalian cell. In contrast, conventional gene therapy vectors require that one disable viral functions, such as expression of viral genes and viral genome replication.

In the present method, the viral particle serves as a "shell" for the delivery of DNA to the mammalian cell. The viral DNA is engineered to contain transcriptional control sequences that are active in a mammalian cell, to allow expression of the gene of interest in the target cell. Conventional recombinant DNA techniques can be used for inserting such sequences. Because the non-mammalian DNA viruses used in the invention are not capable of replicating in mammalian cells, it is not necessary to delete essential viral functions to render them defective. It is preferred, however, that the virus naturally replicate in a eukaryotic species (e.g., an insect, a plant, or a fungus). Examples of viruses that can be engineered to express an exogenous gene in accordance with the invention are listed in Table 1. Preferably, the genome of the virus used in the invention is normally transported to the nucleus in its natural host species because nuclear localization signals function similarly in invertebrate and in mammalian cells. The data summarized below show that, in contrast to conventional wisdom, a non-mammalian DNA virus can (1) infect a wide variety of mammalian cells, and (2) direct expression of an exogenous gene in the cells. In addition, expression of an altered coat protein on the surface of a virus particle enhances the ability of the virus to express an exogenous gene in a mammalian cell.

Established methods for manipulating recombinant viruses may be incorporated into these new methods for expressing an exogenous gene in a mammalian cell. For example, viral genes can be deleted from the virus and supplied in *trans* via packaging lines. Deletion of such genes may be desired in order to (1) suppress expression of viral gene products that may provoke an immune response, (2) provide additional space in the viral vector, or (3) provide additional levels of safety in maintaining the virus in a cell.

This invention involves the expression of an altered coat protein(s) on the surface of virus particle to enhance the ability of a non-mammalian DNA virus to infect a mammalian cell and express an exogenous gene in the mammalian cell. Conventional molecular biology techniques and criteria can be used for identifying and expressing on the virus a polypeptide that binds a mammalian cell. Typically, a gene encoding the altered coat protein is

operably linked to a non-mammalian-active promoter, and is expressed from the viral genome. Alternatively, the altered coat protein can be encoded by a sequence contained within a chromosome of a non-mammalian cell in which the virus is propagated. Upon expression of the altered coat protein from the cellular chromosome, the altered coat protein is packaged along with the non-mammalian DNA virus. In yet another alternative method, the altered coat protein can be expressed from the genome of a second virus that co-infects the non-mammalian cell in which the non-mammalian DNA virus is propagated. Thus, upon co-infection and expression of the altered coat protein from the genome of the second virus, the altered coat protein is packaged along with the non-mammalian DNA virus. Regardless of the method used to express the altered coat protein, the non-mammalian DNA virus is maintained under conditions such that the altered coat protein is expressed on the surface of the virus particle. To this end, conventional methods for propagating viruses (further discussed below) in non-mammalian cells can be used. If desired, expression of the altered coat protein on the surface of a virus particle can be confirmed using conventional techniques, such as immunoblotting, immunofluorescence, and the like.

Conventional molecular biology techniques can be used to produce a suitable fusion protein that is used as the altered coat protein. For example, where a baculovirus is used as the non-mammalian DNA virus, a wide variety of fusion proteins can be made employing the baculovirus coat protein gp64 (Whitford et al., 1989, J. Virol. 63:1393-1399 and Ayres et al., 1994, Virology 202:586-605). The baculovirus expression vector pAcSurf-2 provides a gp64 gene having a multiple cloning site positioned in-phase between the gp64 signal sequence and the sequence encoding the mature glycoprotein (Boublik et al., 1995, Biotechnology 13:1079-1084). Sequences encoding a polypeptide that binds a mammalian cell can readily be inserted into the multiple cloning site of this vector, and expression of the resulting fusion protein is driven by the polyhedrin promoter to which the gp64 sequences are operably linked.

If desired, the viral capsid or envelope can contain, as part of the altered coat protein, or as a separate molecule in addition to the altered coat

protein, a ligand that binds to mammalian cells to facilitate entry. For example, the virus can include as a ligand an asialoglycoprotein that binds to mammalian lectins (e.g., the hepatic asialoglycoprotein receptor), facilitating entry into mammalian cells.

5 Because most promoters of non-mammalian viruses are not active in mammalian cells, the exogenous gene should be operably linked to a promoter that is capable of directing gene transcription in a mammalian cell (i.e., a "mammalian-active" promoter). Examples of suitable promoters include the RSV LTR, the SV40 early promoter, CMV IE promoters (e.g., the human
10 CMV IE1 promoter), the adenovirus major late promoter, and the Hepatitis B viral promoter. Other suitable "mammalian-active" promoters include "mammalian promoters," i.e., sequences corresponding to promoters that naturally occur in, and drive gene expression in, mammalian cells. Often, "mammalian promoters" are also cell-type-specific, stage-specific, or tissue-
15 specific in their ability to direct transcription of a gene, and such promoters can be used advantageously in the invention as a means for controlling expression of the exogenous gene. For example, several liver-specific promoters, such as the albumin promoter/enhancer, have been described and can be used to achieve liver-specific expression of the exogenous gene (see, e.g., Shen et al., 1989,
20 DNA 8:101-108; Tan et al., 1991, Dev. Biol. 146:24-37; McGrane et al., 1992, TIBS 17:40-44; Jones et al., J. Biol. Chem. 265:14684-14690; and Shimada et al., 1991, FEBS Letters 279:198-200). Where the invention is used to treat a hepatocellular carcinoma, an α -fetoprotein promoter is particularly useful. This promoter is normally active only in fetal tissue; however, it is also active in
25 liver tumor cells (Huber et al., 1991, Proc. Natl. Acad. Sci. 88:8039-8043). Accordingly, an α -fetoprotein promoter can be used to target expression of a liver-cancer therapeutic to liver tumor cells.

 If desired, the virus genome can be engineered to carry an origin of replication in order to facilitate persistence of the exogenous gene in the
30 mammalian cell. Origins of replication derived from mammalian cells (i.e., "mammalian origins of replication," have been identified (Burhans et al., 1994, Science 263:639-640). Other origins of replication that function in mammals

(i.e., "mammalian-active" origins, e.g., the Epstein-Barr Virus *oriP*) can also facilitate maintenance of expression in the presence of appropriate *trans*-acting factors (e.g., EBNA-1). If desired, the virus can be engineered to express more than one exogenous gene (e.g., the virus can be engineered to express both
5 OTC and AS) or more than one altered coat protein.

Descriptions of several viruses used in the examples described below now follow. These examples are provided for illustrative purposes, and are not meant to limit the scope of invention.

EXAMPLES OF TRANSFER PLASMIDS

10 Construction of the pZ4 Transfer Plasmid: Genetic manipulation of a baculovirus for use in the invention can be accomplished with commonly-known recombination techniques originally developed for expressing proteins in baculovirus (see, e.g., O'Reilly et al., 1992, *In: Baculovirus expression vectors*, W. H. Freeman, New York). In this example, an AcMNPV was constructed by
15 interrupting the polyhedrin gene of the virus with a cassette that directs expression of a reporter gene. The reporter gene cassette included DNA sequences corresponding to the Rous Sarcoma Virus (RSV) promoter operably linked to the *E. coli lacZ* gene (Fig. 1). The reporter gene cassette also included sequences encoding Simian Virus 40 (SV40) RNA splicing and
20 polyadenylation signals.

The RSV-lacZ AcMNPV transfer plasmid used in several examples set forth below is named Z4 and was constructed as follows. An 847 bp fragment of pRSVPL9 including the SV40 RNA splicing signal and polyadenylation signal was excised using *Bgl*II and *Bam*HI. Plasmid pRSVPL9 was derived
25 from pRSVglobin (Gorman et al., Science 221:551-553) by digesting pRSVglobin with *Bgl*II, adding a *Hind*III linker, and then cleaving the DNA with *Hind*III. A double-stranded polylinker made by hybridization of the oligonucleotides 5'AGCTGTCGACTCGAGGTACCAGATCTCTAGA3' (SEQ ID NO: 1) and 5'AGCTTCTAGAGATCTGGTACCTCGAGTCGAC3' (SEQ
30 ID NO: 2) was ligated to the 4240 bp fragment having the RSV promoter and SV40 splicing and polyadenylation signals. The resulting plasmid has the

polylinker in place of the globin sequences. The SV40 sequence of pRSVPL9 was cloned into the *Bam*HI site of pVL1392 (Invitrogen and Pharmingen) using standard techniques. The resulting intermediate plasmid was named pVL/SV40. An RSV-lacZ cassette was excised from pRSVlacZII (Lin et al., 1991, Biotechniques 11:344-348, and 350-351) with *Bgl*II and *Spe*I and inserted into the *Bgl*II and *Xba*I sites of pVL/SV40.

The AcMNPV RSV-lacZ virus, termed Z4, was prepared by homologous recombination of the Z4 transfer plasmid with linearized AcMNPV DNA. The AcMNPV virus used to prepare this DNA was AcV-EPA (Hartig et al., 1992, J. Virol. Methods 38:61-70).

Construction of the pZ5 Transfer Plasmid: Certain non-mammalian viruses (e.g., baculoviruses) may be occluded in a protein inclusion body (i.e., occluded-derived viruses (ODV)), or they may exist in a plasma membrane budded form. Where an occluded virus is used in the invention, the virus may first be liberated from the protein inclusion body, if desired. Conventional methods employing alkali may be used to release the virus (O'Reilly et al., 1992, *In: Baculovirus expression vectors*, W. H. Freeman, New York). An occluded, alkali-liberated baculovirus may be taken up by a cell more readily than is the non-occluded budded virus (Volkman and Goldsmith, 1983, Appl. and Environ. Microbiol. 45:1085-1093). To construct the pZ5 transfer plasmid (Fig. 2), for using an occluded virus in the invention, the RSV-lacZ cassette was excised from the pZ4 transfer plasmid using *Bgl*II and *Bam*HI and then inserted into the *Bgl*II site of pAcUW1 (Weyer et al., 1990, J. Gen. Virol. 71:1525-1534).

Construction of the pZ-EBV#1 Transfer Plasmid: The non-mammalian DNA viruses used in the invention may be engineered to permit episomal replication of the virus in the mammalian cell. Such a virus would persist longer, thereby optimizing methods for long-term expression of an exogenous gene in a cell. An example of such a replicating virus is pZ-EBV#1 (Fig. 3), which was constructed as follows. The EBV *oriP* and EBNA-1 region was excised from pREP9 (Invitrogen) using *Eco*RI and *Xba*I and then inserted into the baculoviral transfer plasmid pBacPAK9 (Clontech) at its *Eco*RI and *Xba*I

sites, yielding pEBVBP9. The RSV-lacZ cassette was excised from transfer plasmid Z4 with *Bgl*II and *Bam*HI and then inserted into the *Bam*HI site of pEBVBP9 to yield the plasmid pZ-EBV#1.

5 Construction of pZ4loxP: The Z4loxP viral genome is a substrate for recombination with bacteriophage P1 cre recombinase. This virus can be used to insert gene cassettes bearing a loxP site into the virus using standard procedures (Patel et al., 1992, Nucl. Acids Res. 20:97-104). A variation of this insertion system may be engineered so that the viral sequences are excised from the remaining gene expression sequences. For example, an auto-excising transfer plasmid may be constructed (Figs. 4A - 4B) to express an exogenous gene in a mammalian cell. This plasmid contains loxP sequences which facilitate excision of the baculoviral sequences. The pZ4loxP transfer plasmid was constructed by inserting a synthetic loxP site into the pZ4 transfer plasmid. Two loxP oligonucleotides were synthesized and annealed to each other. The oligonucleotides were:

10 5'GATCTGACCTAATAACTTCGTATAGCATACATTATACGAAGTTATAT
TAAGG3' (SEQ ID NO: 3) and
5'GATCCCTTAATATAACTTCGTATAATGTATGCTATACGAAGTTATTA
GGTCA3' (SEQ ID NO:4). The oligonucleotides were annealed by heating
20 them to 80° C in the presence of 0.25 M NaCl and then allowing the mixture to cool slowly to room temperature before use in the ligation reactions. The annealed oligonucleotides were then ligated to the pZ4 transfer plasmid that had been digested with *Bgl*II. The ligations and analysis of the resulting clones were performed with standard cloning techniques. Recombinant Z4loxP
25 baculovirus was then generated with conventional methods for recombination into linear baculoviral DNA.

30 Construction of pBV-AVneo, an AAV Chimera Transfer Plasmid: A baculovirus genome that is capable of integrating into a chromosome of the host cell can also be used in the invention. Such an integrated virus may persist in the cell longer than a non-integrated virus. Accordingly, methods of gene expression involving such viruses may obviate the need for repeated administration of the virus to the cell, thereby decreasing the likelihood of

mounting an immune response to the virus. The transfer plasmid pBV-AVneo (Fig. 5) includes the inverted terminal repeats of an Adeno-associated virus (AAV). This transfer plasmid was constructed by excising the *neo* gene, which encodes G418-resistance, as a *Bgl*II-*Bam*HI fragment from pFasV.neo and
5 inserting the fragment into the *Bam*HI site of pAVgal in place of the *lacZ* gene. Plasmid pAVgal was constructed by replacing the rep and cap coding sequences of AAV with a CMV promoter and a *lacZ* gene. The resulting intermediate fragment, termed pAV.neo, was digested with *Pvu*I. The large *Pvu*I fragment, which has the CMV promoter driving expression of the *neo* gene, flanked by
10 the AAV ITRs, then was inserted into the *Pac*I site of pBacPAK9. If desired, a suitable promoter operably linked to an AAV rep gene may be inserted into this construct (e.g., between the AAV ITR and the polyhedrin promoter) to facilitate excision and recombination into the genome. Examples of rep genes that may be inserted into this construct include rep40, rep52, rep68, and rep78.

15 Construction of the pCMV-BV Transfer Plasmid: The human cytomegalovirus immediate early promoter, a 758 bp *Hind*III-*Xba*I fragment, was excised from pCMV-EBNA (Invitrogen) at *Hind*III, *Bam*HI and inserted into the *Hind*III sites of pBluescript (SKII⁺), yielding plasmid pCMV-SKII⁺. The promoter was then excised from CMV-SKII⁺ at the *Xho*I, *Bam*HI sites and
20 inserted into the *Xho*I, *Bgl*II sites of pSV/BV, yielding plasmid pCMV-BV (Fig. 6). pSV/BV is a modified version of the baculovirus transfer plasmid pBacPAK9 (Clontech), containing an altered polylinker and SV40 splice and polyadenylation signals. pSV/BV was constructed by restriction of pBacPAK9 with *Not*I, treatment with T4 DNA polymerase to create blunt ends, and self-
25 ligation to remove the *Not*I site. A new *Not*I site was then added by ligation of the linker pGCGGCCGC into the *Sma*I site. Finally, SV40 splice and polyadenylation sequences were added by digestion of pRSVPL with *Bgl*II-*Bam*HI, and insertion of the 847 bp fragment into the *Bam*HI site of the modified BacPAK9, yielding pSV/BV.

30 Construction of the pCMVZ-BV Transfer Plasmid: pCMVZ-BV (Fig. 7) was constructed by restriction of pCMV-BV with *Not*I and ligation insertion

of a 3 kb lacZ fragment. The lacZ fragment was prepared by restriction of pAlb-Gal with *NotI*.

Construction of the pAct-BV Transfer Plasmid: The 345 bp rat β -actin promoter was excised from pINA (Morgenstern, JP, 1989, Ph.D. Thesis, University College, London, UK) at *Bgl*II, *Bam*HI and inserted into the *Bgl*II site of pSV/BV, yielding pAct-BV (Fig. 8).

Construction of the pAZ-BV Transfer Plasmid: pAZ-BV (Fig. 9) was constructed by restriction of pAct-BV with *NotI* and ligation insertion of a 3 kb lacZ fragment. The lacZ fragment was prepared by restriction of pAlb-Gal with *NotI*.

Construction of the pIE45-BV Transfer Plasmid: pIE45-BV (Fig. 10) was constructed by restriction of pHSVPrPUC (Neve et al., 1997, Neuroscience 79:435-447) with *Sph*I, followed by treatment with T4 DNA polymerase in the presence of nucleotide triphosphates to create blunt ends. *Pst*I linkers (New England Biolabs, Catalog #1024, pGCTGCAGC) were then added by treatment with T4 DNA ligase, the fragment of approximately 850 bp was subjected to digestion with *Pst*I, and cloned into the *Pst*I site of pSV/BV.

Construction of the pNSE4-BV Transfer Plasmid: pNSE4-BV (Fig. 11) was constructed by restriction of pNSE4 (see, e.g., Quon et al., 1991, Nature 352::239-241 and Forss-Petter et al., 1990, Neuron 5:187-197) with *Sa*I and *Eco*RI, followed by ligation into the *Xho*I and *Eco*RI sites of pSV/BV.

Construction of the pTH/SV40/BP9 Transfer Plasmid: pTH/SV40/BP9 (Fig. 12) was constructed by restriction of pTH4.8 Thdno (Banerjee et al., 1992, J. Neuroscience 12:4460-4467) with *Eco*RI and *NotI*, and ligation of the 4.0 kb promoter fragment into pSV/BV, which was also digested with *Eco*RI and *NotI*.

Construction of the pTH-Lac/BP9 Transfer Plasmid: pThlac (Fig. 13) was constructed by restriction of pALB-Gal with *Not* I and isolation of the 3 kb lacZ fragment, which was then ligated into pTH/SV40/BP9 which was also restricted with *Not* I using T4 DNA ligase.

Propagation of Viruses: Conventional methods can be used to propagate the viruses used in the invention (see, e.g., Burleson, et al., 1992,

Virology: A Laboratory Manual, Academic Press, Inc., San Diego, CA and Mahy, *ed.*, 1985, Virology: A Practical Approach, IRL Press, Oxford, UK). Conventional conditions for propagating viruses are suitable for allowing expression of an altered coat protein on the surface of a virus particle used in the invention. For example, the baculoviruses used in the experiments described below were plaque purified and amplified according to standard procedures (see, e.g., O'Reilly et al. *infra* and Summers and Smith, 1987, A Manual of Methods for Baculovirus Vectors and Insect Cell Culture Procedures, Texas Agricultural Experiment Station Bulletin No. 1555, College Station, Texas). AcMNPV and Sf21 cells were propagated by spinner culture in Hinks TNM-FH media (JRH Biosciences) containing 10% fetal bovine serum (FBS) and 0.1% PLURONIC F-68™. Amplified virus can be concentrated by ultracentrifugation in an SW28 rotor (24,000 rpm, 75 minutes) with a 27% (w/v) sucrose cushion in 5 mM NaCl, 10 mM Tris pH 7.5, and 10 mM EDTA. The viral pellet is then resuspended in phosphate-buffered saline (PBS) and sterilized by passage through a 0.45 µm filter (Nalgene). If desired, the virus may be resuspended by sonication in a cup sonicator. AcMNPV was titered by plaque assay on Sf21 insect cells.

EXAMPLES OF EXOGENOUS GENE EXPRESSION

Because non-mammalian DNA viruses were long thought not to be capable of infecting and directing gene expression in mammalian cells, Part A of the Examples below provides evidence that a non-mammalian DNA virus (e.g., a baculovirus) can, in fact, be used to express an exogenous gene in a mammalian cell. Although the examples described in Part A employ viruses in which the protein coat was not altered, these examples provide support for the assertions that a non-mammalian DNA virus can be used to express an exogenous gene in a mammalian cell. In addition, these examples provide guidance for practicing the invention with a virus having an altered coat protein.

The examples in Part B, below, utilize non-mammalian DNA viruses that have an altered coat protein. Because the presence of the altered coat

protein is the only significant difference between the viruses of the invention and the viruses that lack an altered coat protein, these examples demonstrate that the expression of the altered coat protein enhances the ability of a non-mammalian DNA virus to express an exogenous gene in a mammalian cell.

5 Accordingly, in each of the methods described below (e.g., *in vivo* expression of an exogenous gene), the viruses having an altered coat protein are expected to be superior to the viruses lacking the altered coat protein.

**PART A: A NON-MAMMALIAN DNA VIRUS CAN BE
USED TO EXPRESS AN EXOGENOUS GENE IN A MAMMALIAN CELL**

10 **I. Examples of Expression of an Exogenous Gene in Mammalian Cells *In Vitro***

Nearly all mammalian cells are potential targets of non-mammalian viruses, and any cultured or primary cell can rapidly be tested. In the following example, the ability of the Z4 baculovirus to infect 19 different types of cells was tested. In this example, the baculovirus was the Z4 virus, prepared

15 by homologous recombination of the Z4 transfer plasmid with linearized AcMNPV DNA. The tested cells were HepG2, Sk-Hep-1, NIH3T3, NIH3T3 cells expressing a cell-surface asialoglycoprotein receptor, HeLa, CHO/dhfr, 293, COS, Ramos, Jurkat, HL60, K-562, C₂C₁₂ myoblasts, C₂C₁₂ myotubes, primary human muscle myoblasts, Hep3B cells, FTO2B cells, Hepa1-6 cells,

20 and nerve growth factor-differentiated PC12 cells.

Growth of Cells: Conventional tissue culture methods can be used to grow mammalian cells to be infected (Freshney, 1987, Culture of Animal Cells: A Manual of Basic Techniques, 2nd ed., Alan R. Liss, Inc. New York, NY). These cells were grown and infected as is described above. The cells were

25 grown as follows. HepG2 and Sk-Hep-1 cells were cultured in minimal essential medium as modified by Eagle (EMEM) containing 10% FBS. NIH3T3, HeLa, 293, and COS cells were cultured in DMEM containing 10% FBS. CHO/dhfr cells were cultured in MEM alpha containing 10% FBS. Ramos, Jurkat, HL60, and K-562 cells were cultured in RPMI 1640 medium

30 containing 10% FBS. HL60 cells were induced to differentiate by culture in

the same medium containing 0.5% dimethyl sulfoxide and 1 μ M retinoic acid (Sigma). C₂C₁₂ myoblasts were propagated in DMEM containing 20% FBS and differentiated to myotubes during culture in DMEM containing 10% horse serum. PC12 cells were propagated in DMEM containing 5% FBS and 10% horse serum, and were induced to differentiate during culture in DMEM containing 10% FBS, 5% horse serum, and 100 ng/ml nerve growth factor. All cells were seeded one day prior to infection with AcMNPV, and multiplicities of infection were calculated assuming a doubling in cell number during this time. The C₂C₁₂ and PC12 cells may have increased in cell number during differentiation, and therefore reflect a somewhat lower moi.

In vitro Infection of Cells: *In vitro* infection of mammalian cells with a virus can be accomplished by allowing the virus to adsorb onto the cells for 0.1 to 6 hours; preferably, adsorption proceeds for 1 to 2 hours. Generally, a multiplicity of infection of 0.1 to 1,000 is suitable; preferably, the moi is 100 to 500. For relatively refractory cells, a moi of 100 to 1,000 is preferable. For the viruses used in the invention, the titer may be determined with conventional methods which employ the non-mammalian cells that the virus naturally infects. If desired, the mammalian cell to be infected may be maintained on a matrix that contains collagen (e.g., rat tail Type I collagen). Based on cell counting after culture and infection of cells on collagen-coated plates and comparison with cells grown on a conventional EHS matrix, I have found that a collagen matrix increases the susceptibility of cells (e.g., liver cells) to infection by a non-mammalian virus by 10 to 100 fold, relative to a conventional EHS matrix. Commercially-available plates containing a collagen matrix are available (e.g., BIO-COATTM plates, Collaborative Research), and rat tail collagen is also commercially available (Sigma Chemical and Collaborative Research).

In the *in vitro* assays described below, standard conditions for infection utilized 2 x 10⁶ cells and RSV-lacZ AcMNPV at a moi of 15. Adherent cell lines were seeded one day prior to infection. Cells were exposed to virus in 2 ml of medium for 90 minutes, and then the virus-containing medium was removed and replaced with fresh medium. Mock-infected cells were treated with 2 ml medium lacking the viral inoculum.

Detection of Infection and Gene Expression: Delivery of a virus to a cell and expression of the exogenous gene can be monitored using standard techniques. For example, delivery of a virus (e.g., AcMNPV) to a cell can be measured by detecting viral DNA or RNA (e.g., by Southern or Northern blotting, slot or dot blotting, or *in situ* hybridization, with or without amplification by PCR). Suitable probes that hybridize to nucleic acids of the virus, regulatory sequences (e.g., the promoter), or the exogenous gene can be conveniently prepared by one skilled in the art of molecular biology. Where the invention is used to express an exogenous gene in a cell *in vivo*, delivery of the virus to the cell can be detected by obtaining the cell in a biopsy. For example, where the invention is used to express a gene in a liver cell(s), a liver biopsy can be performed, and conventional methods can be used to detect the virus in a cell of the liver.

Expression of an exogenous gene in a cell of a mammal can also be followed by assaying a cell or fluid (e.g., serum) obtained from the mammal for RNA or protein corresponding to the gene. Detection techniques commonly used by molecular biologists (e.g., Northern or Western blotting, *in situ* hybridization, slot or dot blotting, PCR amplification, SDS-PAGE, immunostaining, RIA, and ELISA) can be used to measure gene expression. If desired, a reporter gene (e.g., *lacZ*) can be used to measure the ability of a particular baculovirus to target gene expression to certain tissues or cells. Examination of tissue can involve: (a) snap-freezing the tissue in isopentane chilled with liquid nitrogen; (b) mounting the tissue on cork using O.C.T. and freezing; (c) cutting the tissue on a cryostat into 10 μ m sections; (d) drying the sections and treating them with 4% paraformaldehyde in PBS, followed by rinsing in PBS; (e) staining the tissue with X-gal (0.5 mg/ml)/ ferrocyanide (35 mM)/ ferricyanide (35 mM) in PBS; and (f) analyzing the tissue by microscopy.

To measure expression of the reporter gene in the infected cells, colorimetric assays of β -galactosidase enzymatic activity were performed with standard methods (Norton et al., 1985, Molecular & Cellular Biology 5:281-290). Other conventional methods for measuring β -galactosidase activity could be used in lieu of the methods employed in this example. Cell extracts were

prepared at one day post-infection. Cell monolayers were rinsed three times with PBS, scraped from the dish, and collected by low-speed centrifugation. The cell pellets were resuspended in 25 mM Tris pH 7.4/0.1 mM EDTA and then subjected to three cycles of freezing in liquid nitrogen and thawing in a 37°C water bath. The extracts were then clarified by centrifugation at 14,000 x g for 5 minutes. Standard conditions for assaying β -galactosidase activity utilized 0.1 ml of cell extract, 0.8 ml of PM-2 buffer, and 0.2 ml of *o*-nitrophenyl- α -D-galactopyranoside (4 mg/ml) in PM-2 buffer for 10 minutes at 37°C (Norton et al., 1985, Mol. & Cell. Biol. 5:281-290). The reaction was stopped by the addition of 0.5 ml of 1 M sodium carbonate. The amount of substrate hydrolyzed was detected spectrophotometrically at 420 nm, and β -galactosidase enzymatic activity was calculated with conventional methods (Norton et al., 1985, Mol. & Cell. Biol. 5:281-290). The assay was verified to be linear with respect to extract concentration and time. Extract protein concentrations were determined using the Coomassie Plus protein assay (Pierce) with bovine serum albumin as a standard, and the level of β -galactosidase activity was expressed as units of β -galactosidase activity per mg of protein. Other standard protein assays can be used, if desired.

For histochemical staining of β -galactosidase activity, cells were fixed in 2% (w/v) formaldehyde-0.2% (v/v) paraformaldehyde in PBS for 5 minutes. After several rinses with PBS, the cells were stained by the addition of 0.5 mg/ml of X-gal (BRL) in PBS for 2-4 hours at 37°C.

Assay of 19 Mammalian Cell Types: The following 19 examples illustrate that expression of an exogenous gene can be detected in 14 of the 19 mammalian cell types that were tested. These assays employed two different tests of β -galactosidase activity. By X-gal staining, the more sensitive assay, exogenous gene expression was detected in 14 of the 19 mammalian cell types. Using an ONPG assay of cell extracts, which is a less sensitive assay, three of the cell lines (HepG2, 293, and PC12) showed statistically significant ($P < 0.05$, Student's t-test) higher β -galactosidase activity after exposure to the virus (Table 3). The human liver tumor line HepG2 exposed to the RSV-*lacZ* baculovirus expressed greater than 80-fold higher levels of β -galactosidase than

did mock-infected controls. The adenovirus-transformed human embryonal kidney cell line 293 expressed the *lacZ* reporter gene at a level of about four-fold over background. In addition, PC12 cells, which were differentiated to a neuronal-like phenotype with nerve growth factor, exhibited about two-fold higher β -galactosidase levels after infection with the RSV-*lacZ* baculovirus. This difference was statistically significant ($P=0.019$).

TABLE 3. BACULOVIRUS-MEDIATED EXPRESSION OF AN RSV-LACZ REPORTER GENE IN MAMMALIAN CELL LINES.

		β -galactosidase activity (units/mg) Mean \pm SD	
	Cell Line	Mock Infected	RSV- <i>lacZ</i> Virus
	HepG2	0.030 \pm 0.004	2.628 \pm 0.729
	Sk-Hep-1	0.019 \pm 0.003	0.019 \pm 0.004
	NIH3T3	0.026 \pm 0.003	0.023 \pm 0.005
15	HeLa	0.034 \pm 0.009	0.036 \pm 0.005
	CHO/dhfr-	0.020 \pm 0.002	0.026 \pm 0.005
	293	0.092 \pm 0.014	0.384 \pm 0.024
	COS	0.029 \pm 0.002	0.032 \pm 0.007
	Ramos	0.008 \pm 0.002	0.011 \pm 0.004
20	Jurkat	0.012 \pm 0.004	0.007 \pm 0.001
	HL60	0.042 \pm 0.039	0.014 \pm 0.015
	K-562	0.018 \pm 0.006	0.017 \pm 0.002
	C ₂ C ₁₂ myoblast	0.015 \pm 0.001	0.014 \pm 0.003
	C ₂ C ₁₂ myotube	0.049 \pm 0.011	0.042 \pm 0.004
25	PC12 (+NGF)	0.019 \pm 0.005	0.033 \pm 0.004

By histochemical staining, a more sensitive assay, β -galactosidase activity was detected in 14 of the 19 cell lines exposed to virus. Thus, certain of the cell lines that did not yield statistically significantly higher levels of β -galactosidase, as measured in extracts, were, in fact, able to express β -galactosidase at low, but reproducible, frequencies, as detected by the more sensitive X-gal staining procedure. This frequency could be increased by using

higher multiplicities of infection such that cells that, at a low moi appear not to express the gene, stain blue at a higher moi. Examples of cell lines that could be transfected in this manner include SK-Hep-1, NIH3T3, HeLa, CHO/dhfr^r, 293, Cos, and C₂C₁₂ cells. In addition, β -galactosidase activity was detected in
5 primary human muscle myoblasts that were exposed to virus. This finding indicates that baculovirus was able to mediate gene transfer both to primary cells and the corresponding established cell line (C₂C₁₂), indicating that expression of the exogenous gene in an established cell line has predictive value for the results obtained with primary cells.

10 β -galactosidase activity was also detected in Hep3B cells treated with the virus; the level of expression in these cells was nearly equivalent to the level detected with HepG2 cells. In addition, β -galactosidase activity was found in FTO2B (rat hepatoma) cells and Hepa1-6 (human hepatoma) cells exposed to virus. β -galactosidase activity was also detected in NIH3T3 cells that were
15 engineered to express the asialoglycoprotein receptor on the cell surface. These cells expressed approximately two times the level of β -galactosidase as did normal NIH3T3 cells. This observation suggests that an asialoglycoprotein receptor may be used to increase susceptibility to viral-mediated gene transfer.

At the moi employed, the Ramos, Jurkat, HL60, and K-562 cell lines did
20 not express statistically significant levels of β -galactosidase, as revealed by β -galactosidase enzyme assays after infection. Based on the results with other mammalian cell lines, it is expected that β -galactosidase activity would be detected in these apparently refractory cell lines when a higher dose (i.e., moi) of virus or longer adsorption time period is utilized.

25 Even when exposure of cells to the virus results in expression of the exogenous gene in a relatively low percentage of the cells (*in vitro* or *in vivo*), the invention can be used to identify or confirm the cell- or tissue-type specificity of the promoter that drives expression of the exogenous gene (e.g., a reporter gene such as a chloramphenicol acetyltransferase gene, an alkaline
30 phosphatase gene, a luciferase gene, or a green fluorescent protein gene). Once identified, such a promoter may be employed in any of the conventional methods of gene expression. Similarly, only relatively low levels of expression

are necessary for provoking an immune response (i.e., produce antibodies) in a mammal against the heterologous gene product. Thus, the gene expression method of the invention can be used in the preparation of antibodies against a preferred heterologous antigen by expressing the antigen in a cell of a mammal.

5 Such antibodies may be used *inter alia* to purify the heterologous antigen. The gene expression method may also be used to elicit an immunoprotective response in a mammal (i.e., be used as a vaccine) against a heterologous antigen. In addition, the invention can be used to make a permanent cell line from a cell in which the virus mediated expression of a cell-immortalizing

10 sequence (e.g., SV40 T antigen).

Histochemical staining using X-gal provided a highly sensitive method for detecting β -galactosidase expression in cells exposed to the modified AcMNPV. When HepG2 cells were exposed to the modified AcMNPV at a moi of 15, about 5-10% of the cells stained with X-gal (Fig. 14A). At a

15 multiplicity of infection (moi) of 125, about 25-50% of the cells were stained (Fig. 14B). No adverse effects of exposure to the virus, such as nuclear swelling, were observed. These data demonstrate that the modified AcMNPV is highly effective at gene transfer into HepG2 cells when a sufficient dose of virus is used. When the Sk-Hep-1 line was exposed to virus at a moi of 15, no

20 stained cells were observed (data not shown). While the majority of Sk-Hep-1 cells that were exposed to virus at a moi of 125, did not stain blue (Fig. 14C), a few cells were found that stained darkly after treatment with this higher doses of virus (Fig. 14D). These data indicate that cells that appear to be refractory to the virus at a relatively low moi can, in fact, be infected, and express the

25 exogenous gene, at a higher moi. Stained cells were not found in mock-infected cultures (data not shown). The frequency of stained cells in the Sk-Hep-1 cell line was estimated to be 2,000-4,000 fold less than in HepG2 cells after exposure to equivalent doses of the modified virus, as determined by cell counting. Thus, the cell type-specificity demonstrated by the modified

30 AcMNPV is relative rather than absolute. These data also indicate that, where a mixture of cells is contacted with the virus (*in vitro* or *in vivo*), the dosage of

the virus can be adjusted to target the virus to the cells that are infected at a lower moi.

Expression in Primary Cultures of Rat Hepatocytes: This example illustrates that a non-mammalian DNA virus can also be used to express an exogenous gene at high levels in primary cultures of rat hepatocytes. In this experiment, freshly prepared rat hepatocytes were plated onto dishes coated with rat tail collagen as previously described (Rana et al., 1994, Mol. Cell. Biol. 14:5858-5869). After 24 hours, the cells were fed with fresh medium containing RSV-*lacZ* baculovirus at a multiplicity of infection of approximately 430. After an additional 24 hours, the cells were fixed and stained with X-gal. Over 70% of the cells were stained blue, indicating that they have taken up and expressed the RSV-*lacZ* cassette (Fig. 15). The frequency of expression obtained in this example is higher than the frequency reported with conventional viral vectors used in gene therapy (e.g., retroviral and Herpes Simplex Virus vectors). Mock-infected cultures did not contain any positively-stained cells (data not shown). Other preferred exogenous genes can be used in lieu of the *lacZ* gene. In addition, other primary cells can readily be plated and incubated with a non-mammalian cell in lieu of the primary rat hepatocytes.

Expression in Cortex Cultures: The following two examples illustrate that a non-mammalian DNA virus can be used to express an exogenous gene in cultured neuronal and glial cells. For this example, the Z4 virus was prepared from Sf9 cells grown in Hink's TNM-FH media containing 10% FCS, as described above. The virus was purified by banding on a 20-60% sucrose gradient in phosphate-buffered saline. The titer of the virus employed in the following experiments was 3×10^8 pfu/ml (for virus stock #1) or 2×10^9 pfu/ml (for virus stock # 2), as measured on Sf9 cells. Each virus stock was sonicated prior to use.

For the first example, rat cerebral cortex cultures were prepared from E16 embryonic pups. A 24-well dish was seeded with 300,000 cells/well, and, at 4 days post-plating, the cells were infected by adding varying amounts of virus in serum-containing medium to the wells, as is indicated in Table 4. The virus was allowed to adsorb onto the cells for 24 hours.

TABLE 4. EXPRESSION OF AN EXOGENOUS GENE IN RAT CORTICAL CELLS.

VIRUS	1 μl	2 μl	5 μl	10 μl	50 μl	100 μl
Z4 Stock #1	moi = 1	moi = 2	moi = 5	moi = 10	moi = 50	moi = 100
	no blue cells	no blue cells	~5 blue cells	~20 blue cells	~500 blue cells	~2200 blue cells (~0.75%)
Z4 Stock #2	moi = 6.7	moi = 13.3	moi = 34	moi = 67	moi = 335	moi = 667
	few blue cells	~100 blue cells	~200 blue cells	~450 blue cells	~1000 blue cells	~1300 blue cells
PBS				no blue cells	no blue cells	no blue cells

Expression of the exogenous β -galactosidase gene was measured by counting the number of blue cells after staining the cells with X-gal. Table 4 provides the number of blue cells observed in five fields of the microscope at 10X magnification; each well contained approximately 65 fields. In some wells, the cells at the periphery of the well were preferentially stained.

These data indicate that the exogenous β -galactosidase gene was expressed from the virus in the cultured neuronal cells. In contrast, no blue cells were detected when the cell cultures were mock-infected with PBS. Thus, this non-mammalian virus can be used to express an exogenous gene in neuronal and glial cells, as determined by the detection of blue cells that were, by cell morphology, identified as neurons and glia according to standard criteria.

In the second example, the Z4 baculovirus was used to express an exogenous gene in cultured cortical cells obtained from rat pups at the E20 and P1 stages. The cells from E20 pups were plated in 24-well dishes at 380,000 cells/well. The cells from P1 pups were plated at 300,000 cells/well. The E20 cultures were treated with araC (to inhibit the growth of glia) at 6 days post-plating, and they were infected at 10 days post-plating. The P1 cultures were treated with araC at 2 days post-plating, and they were infected at 6 days post-plating. Samples of each culture were infected with various dilutions of Z4 virus at titer 2×10^9 pfu/ml. To measure the strength of the RSV promoter, the cells were also infected, in separate experiments, with Herpes Simplex Virus

(HSV) expressing the *lacZ* gene under two different promoters. In one case, cells were infected with a HSV in which the *lacZ* gene was placed under the control of an RSV promoter. The titer of this HSV stock was 2×10^7 IU/ml, as measured on PC12 cells with X-gal histochemistry. For comparison, the cells were infected with a HSV in which the *lacZ* gene was placed under control of the HSV IE4/5 promoter. The titer of this virus was 2×10^8 IU/ml, as measured on PC12 cells with X-gal histochemistry. For a negative control, the cells were mock-infected with PBS. Expression of the exogenous *lacZ* gene was measured by counting the number of blue cells obtained upon staining the cells with X-gal.

The non-mammalian Z4 virus of the invention successfully expressed the exogenous *lacZ* gene in cultured cortical cells obtained from rat pups at both the E20 and P1 stages of development. With 1-100 μ l of the Z4 virus, 4.9-10% of the cortical cells at the E20 stage, and 2.1-5.75% of the cortical cells at the P1 stage, were stained blue with X-gal, indicating expression of the exogenous gene in those cells. Of the cells infected with 0.1-5.0 μ l of the HSV RSVlacZ virus, as a positive control, 1.9-3.4% of the E20 cells, and 0.45-4.2% of the P1 cells stained blue with X-gal. When the cells were infected with a 5 μ l sample of HSV expressing *lacZ* from the IE4/5 promoter, nearly 100% of the cells stained blue. When E20 or P1 cortical cells were mock-infected with PBS, as a negative control, no blue cells were detected. These data provide additional evidence that the non-mammalian Z4 baculovirus can be used to express an exogenous gene in cortex cells. These data also indicate that the level of expression obtained with the Z4 virus is comparable to the level of expression obtained with HSV.

Dose-response of Baculovirus-mediated Gene Transfer: The histochemical data presented above indicate that increasing amounts of β -galactosidase are produced after exposure of mammalian cells to increasing amounts of virus. To quantitate the dose-dependence of baculovirus-mediated gene expression, HepG2 cells were exposed to increasing doses of virus and assayed for β -galactosidase enzyme activity. The amount of enzyme produced was linearly related to the inoculum of virus used over a wide range of doses

(Fig. 16). This suggests that entry of each virus particle occurs independently of entry of other virus particles. The maximum dose of virus used in this assay was limited by the titer and volume of the viral stock, and no plateau in the amount of expression was observed using higher doses of virus. Accordingly, these data indicate that, in practicing the invention, one can modulate the level of expression (i.e., the percent of cells in which the exogenous gene is expressed) by adjusting the dosage of virus used.

Time Course of Baculovirus-mediated Gene Transfer: HepG2 cells were exposed to the RSV-*lacZ* virus for 1 hour, after which the cells were harvested at various times and quantitatively assayed for β -galactosidase activity. As is shown in Fig. 17, β -galactosidase activity was detected as early as 6 hours after exposure to the virus, and expression peaked 12-24 hours post-infection. As is expected for an episomal DNA molecule, expression from the RSV-*lacZ* cassette gradually subsided at later time (Fig. 17 and data not shown). *LacZ* expression remained detectable by X-gal staining at 12 days post-transfection in fewer than 1 in 1,000 cells (data not shown). This expression of *LacZ* was not the result of viral spread, because culture supernatants taken from HepG2 cells 10 days post-infection had titers of 10 pfu/ml as determined by plaque assay on Sf21 cells. These data suggest that, where the invention is used in the manufacture of proteins that are purified from HepG2 cells, it may be desirable to isolate the protein from the cell at a time not sooner than 6 hours after infection of the cell. Depending on the half-life of the protein, it may be desirable to isolate the protein shortly after the peak in protein expression (i.e., after approximately 22-26 hours (e.g., approximately 24 hours) post-infection for HepG2 cells). The optimal time period for maximizing isolating the manufactured protein can readily be determined for each protein, virus, and cell.

Expression Occurs *De Novo* in Mammalian Cells: These examples confirm that expression of the exogenous gene occurs *de novo* in mammalian cells. To demonstrate that the detected reporter gene activity in the mammalian cells was not simply the result of β -galactosidase being physically associated with AcMNPV virions as they enter the mammalian cell, several experiments

were performed that demonstrate that the observed expression of the *lacZ* reporter gene was the result of *de novo* synthesis of β -galactosidase. First, the RSV-*lacZ* virus inoculum was assayed for β -galactosidase activity, and the level of β -galactosidase activity was found to be less than 10% of that expressed after infection of HepG2 cells. Second, HepG2 cells were infected with the RSV-*lacZ* virus and then cultured in the presence of the protein synthesis inhibitor cycloheximide. Inclusion of cycloheximide after infection inhibited the accumulation of β -galactosidase enzyme activity by more than 90% (Table 5). Third, HepG2 cells were infected at an equivalent moi with BacPAK6 (Clontech), a baculovirus in which the *lacZ* gene was under control of the viral polyhedrin promoter rather than the RSV promoter (Table 5). The latter virus expresses extremely high levels of β -galactosidase activity in insect cells where the promoter is active (data not shown). In mammalian cells, the viral polyhedrin promoter is inactive, and the virus containing this promoter failed to provide any enzyme activity in mammalian cells (Table 5). In contrast to prior studies of baculovirus interactions with mammalian cells, these data demonstrate that *de novo* synthesis of *lacZ* occurs after baculovirus-mediated gene transfer into a mammalian cell.

TABLE 5. BACULOVIRUS-MEDIATED GENE EXPRESSION OCCURS *DE NOVO*.

Virus	Drug During Infection	Drug Post Infection	β -galactosidase (% of RSV- <i>lacZ</i> , mean \pm SD)
RSV- <i>lacZ</i>	none	none	100 \pm 5.8
none	none	none	3.2 \pm 0.4
RSV- <i>lacZ</i>	none	cycloheximide	10.3 \pm 1.0
BacPAK6	none	none	2.8 \pm 0.4
RSV- <i>lacZ</i>	chloroquine	chloroquine	2.9 \pm 0.1
RSV- <i>lacZ</i>	none	chloroquine	25.1 \pm 6.2

Baculovirus-mediated Gene Transfer is Inhibited by Lysomotropic

Agents: To gain insight into the mechanism by which baculoviruses express an exogenous gene in a mammalian cell, the susceptibility of gene expression to a lysomotropic agent was examined. Like other enveloped viruses, the budded
5 form of AcMNPV normally enters cells via endocytosis, followed by low pH-triggered fusion of the viral envelope with the endosomal membrane, thus allowing escape into the cytoplasm (Blissard et al., 1993, J. Virol. 66:6829-6835; Blissard et al., 1990, Ann. Rev. of Entomol. 35:127-155). To determine whether endosome acidification was necessary for baculovirus-mediated gene
10 transfer into mammalian cells, HepG2 cells were infected with RSV-*lacZ* AcMNPV in the presence of chloroquine, a lysomotropic agent. HepG2 cells were exposed to AcMNPV virus in media containing or lacking inhibitor for 90 minutes, then the virus-containing media were removed and replaced with fresh media containing or lacking inhibitors as listed.

15 At one day post-infection, the cells were harvested and extracts were assayed for β -galactosidase activity and protein content. Each value in the table represents the average of three independent assays, with the amount of β -galactosidase produced by the RSV-*lacZ* AcMNPV virus in the absence of inhibitors assigned a value of 100%. β -galactosidase activity was normalized
20 for protein content of each extract. When 25 μ M chloroquine was continuously present during and after exposure of HepG2 cells to the virus, *de novo* expression of β -galactosidase was completely prevented (Table 5). This suggests that baculovirus-mediated gene transfer is dependent upon endosomal acidification. When chloroquine was added to the cells at 90 minutes after
25 exposure to the virus, only partial inhibition of β -galactosidase expression was observed. Apparently, a portion (\approx 22%) of the viral particles were able to proceed through the endosomal pathway during the 90 minutes of exposure to the virus.

Baculovirus-mediated Gene Transfer is Enhanced by Butyrate: This
30 example illustrates that butyrate enhances the ability of a baculovirus to express an exogenous gene in a mammalian cell. Five transfer plasmids containing different mammalian promoters were created, as diagrammed in Fig. 21. These

vectors were constructed using pSV/BV, a modified version of the baculovirus transfer plasmid pBacPAK9 (Clontech), containing an altered polylinker and SV40 splice and polyadenylation signals. pSV/BV was constructed by restriction of pBacPAK9 with *NotI*, treatment with T4 DNA polymerase to create blunt ends, and self-ligation to remove the *NotI* site. A new *NotI* site was then added by ligation of the linker pGCGGCCGC into the *SmaI* site. Finally, SV40 splice and polyadenylation sequences were added by digestion of a variant of pRSVglobin with *BglII*-*BamHI*, and insertion of the 850 bp fragment into the *BamHI* site of the modified BacPAK9, yielding pSV/BV.

The human cytomegalovirus immediate early promoter, 758 bp *HindIII*-*XbaI* fragment, was excised from pCMV-EBNA (Invitrogen) at *HindIII*, *BamHI* and inserted into the *HindIII*, *BamHI* sites of pBluescript (SKII⁺), yielding plasmid pCMV-SKII⁺. The promoter was then excised from CMV-SK II⁺ at the *XhoI*, *BamHI* sites and inserted into the *XhoI*, *BglII* sites of pSV/BV, yielding plasmid pCMV/BV. The 500 bp mouse phosphoglycerate kinase (PGK) promoter was prepared by cutting pKJ1-neo (Tybulewicz, 1991, Cell 65: 1153-1163) with *EcoRI* and made blunt with T4 DNA polymerase to remove the *EcoRI* site. The resulting pKJ1 plasmid lacking the *EcoRI* site was amplified by pfu polymerase chain reaction using the primers

5'ACCGCGGATCCAATACGACTCACTATAG3' (SEQ ID NO: 5) and
5'CGGAGATCTGGAAGAGGAGAACAGCGCGGCAG3' (SEQ ID NO: 6).

The amplified PGK promoter was then digested with *XhoI* and *BglII* and inserted into the same sites of pSV/BV yielding PKJ1/BV. The 345 bp rat β -actin promoter was excised from pINA (6) at *BglII*, *BamHI* and inserted into the *BglII* site of pSV/BV yielding p β -actin/BV. The 2.3 kb albumin enhancer and 700 bp albumin promoter were excised from pGEMAlbSVPA (Zaret et al., 1988, Proc. Natl. Acad. Sci. 85: 9076-9080) at *NaeI*, *NsiI* and inserted into the *SmaI*, *PstI* sites of pSV/BV. The RSVlacZ transfer plasmid used (also referred to herein as the Z4 virus) is described above. A 3.0 kb Lac Z cassette was inserted into the *NotI* site of all of the plasmids constructed (See Fig. 21).

Recombinant viruses were generated by cotransfection of the baculovirus transfer vectors with linear BP6 viral DNA (Clontech) into Sf21

cells. The recombinant viruses were purified through three rounds of plaque isolation and amplified on Sf21 cells. The amplified viruses were concentrated by ultracentrifugation as described above and titered by a 96-well method on Sf21 insect cells (O'Reilly et al., 1992, Baculovirus Expression Vectors: A Laboratory Manual, W.H. Freeman, New York, NY).

The human hepatocellular carcinoma cell line HepG2 was infected with each recombinant virus at a multiplicity of infection of 100. Two million cells were infected in a final volume of 1 ml Eagle's Minimum Essential Medium in a 60 mm tissue culture dish. The infection was allowed to proceed for two hours, then 4 ml of complete medium was added to the cells. In a second series of HepG2 infections, the conditions of the first infections were repeated with the exception that after the infection had proceeded for 2 hours 25 μ l of sodium butyrate (100 mM) was added to the cells with 1.5 ml complete media. As a control, cells were mock-infected to assess background β -galactosidase enzyme activity. The cell monolayers were collected after 24 hours and prepared for a colorimetric assay (with ONPG) of β -galactosidase enzymatic activity as described above. Hepatocytes were isolated by collagenase perfusion and plated on rat tail collagen as previously described (Boyce et al., 1996, Proc. Natl. Acad. Sci. 93:2348-2352). Assay conditions (time and amount of extract used) were varied to be within the linear range of the assay. The amount of product was determined by spectrophotometry and β -galactosidase enzyme activity was calculated. The Coomassie Plus protein assay (Pierce) was used to determine the protein concentration of the extracts, and results were expressed as units of β -galactosidase normalized to total protein content of the extract. The amount of background activity from the mock-infected cells was subtracted from the total amount of enzyme activity for each of the promoters. Each infection was performed in triplicate, and expressed as the mean average with standard deviation (Table 6).

As shown in Table 6, the incorporation of viral or mammalian cellular promoters into baculoviruses allows for expression of an exogenous gene product in mammalian cells. The CMV promoter led to the highest level of β -galactosidase activity, with the RSV and β -actin promoters producing lower

levels of β -galactosidase activity. At the moi of virus employed in this example, the albumin and PGK promoters showed no activity above background levels in extracts of cells that were not treated with butyrate, although positively stained cells were detected by X-gal staining. The addition of sodium butyrate to the cells after infection led to detectable levels of β -galactosidase expression with all of the promoters tested. After treating cells with sodium butyrate, the CMV promoter showed a five-fold increase in expression of the β -galactosidase reporter gene. The RSV LTR, albumin, pGK1, and β -actin promoters all led to increased gene expression after treatment with butyrate. Without being bound to any particular theory, it is postulated that sodium butyrate increases cellular differentiation and histone acetylation, which increases transcription.

TABLE 6. COMPARISON OF VARIOUS PROMOTER STRENGTHS WITH AND WITHOUT SODIUM BUTYRATE

	Hep G2	Hep G2	Rat Hepatocytes
Promoter	-butyrate	+butyrate	-butyrate
CMV	17 \pm 1.4 ^a	86 \pm 33	18 \pm 1.2
RSV	1.0 \pm 0.1	2.2 \pm 0.1	0.25 \pm 0.11
pGK1	0.0 \pm 0.0	0.02 \pm 0.02	0.64 \pm 0.58
Albumin	0.0 \pm 0.0	0.08 \pm 0.04	0.15 \pm 0.08
β -actin	0.1 \pm 0.01	0.05 \pm 0.02	0.25 \pm 0.07

^a Promoter strength is expressed in Units/mg of β -galactosidase.

Analysis of RNA Expression From Viral Promoters in HepG2 Cells:

One advantage of using a non-mammalian virus to express an exogenous gene in a mammalian cell is that, due to a lack of appropriate host cell factors, the non-mammalian viral promoters may not be active in the mammalian cell. To determine whether AcMNPV viral gene are expressed in HepG2 cells, the viral RNA was analyzed. In these experiments, HepG2 cells were infected with the Z4 virus at a moi of approximately 30. At 18 hours post-infection, the cells

were harvested, and total cellular RNA was extracted from the cells. The total cellular RNA was analyzed by Northern blotting for expression of viral genes. The probe included a 1.7 kbp *PacI-SalI* fragment from pAcUW1 (Pharmingen) which contains the viral late gene, p74, as well as the very late
5 (hyperexpressed) gene, p10. Total cellular RNA from Z4-infected Sf9 insect cells was employed as a positive control. While extremely strong signals were detected for p10 and p74 for the control insect cells, no signal was observed for Z4-infected HepG2 cells or uninfected control cells.

Additional experiments that used reverse transcriptase-PCR (RT-PCR), a
10 highly sensitive method, provided further evidence that the majority of viral genes are not transcribed in the mammalian HepG2 cells. RT-PCR analysis was performed with RNA prepared from Z4-infected HepG2, uninfected HepG2, or infected Sf9 cells at 6 or 24 hours post-infection. HepG2 cells were infected at a moi of 10 or 100. At 6 hours post-infection, no RT-PCR product
15 was observed from the viral p39, ETL, LEF1, IE1, or IE-N genes at either dose of virus in Z4-infected HepG2 cells. In contrast, RT-PCR products were readily detected in Z4-infected Sf9 cells. At 24 hours post-infection, no expression of these gene was detected in HepG2 cells infected at a moi of 10. At 24 hours post-infection, no expression of the viral p39, ETL, or LEF1 genes
20 was observed in HepG2 cells infected at an moi of 100. However, at this high doses of virus, low levels of expression from the viral IE1 and IE-N genes was observed. The low level of expression detected at an moi of 100 was nonetheless significantly lower than the level of expression in insect cells.

Expression of these genes may result from recognition of the viral TATA
25 box by mammalian transcription factors (i.e., transcription of the immediate early genes by RNA polymerase II (see, e.g., Hoopes and Rorhman, 1991, Proc. Natl. Acad. Sci. 88:4513-4517). In contrast to the immediate early genes, the late or very late viral genes are transcribed by a virally-encoded RNA polymerase that, instead of requiring a TATA box, initiates transcription at a
30 TAAG motif (O'Reilly et al., *supra*). Accordingly, expression of the viral late or very late genes is naturally blocked in mammalian cells. If desired,

expression of the immediate early genes can be blocked by deleting those genes, using conventional methods.

While certain viruses have an intrinsic ability to infect liver cells, infection of liver cells by other viruses may be facilitated by a cellular receptor, such as a cell-surface asialoglycoprotein receptor (ASGP-R). HepG2 cells differ from Sk-Hep-1 human hepatocytes and NIH3T3 mouse fibroblast cells by the presence of ASGP-R on the cell surface. In certain of the above experiments, β -galactosidase was expressed in fewer Sk-Hep-1 cells (Fig. 14B) or NIH3T3 cells than HepG2 cells. The *lacZ* gene was expressed in HepG2 cells at a frequency estimated as greater than 1,000 fold more than that in Sk-Hep-1 cells, based on quantitative counts of X-gal stained cells. Normal hepatocytes have 100,000 to 500,000 ASGP-R, with each receptor internalizing up to 200 ligands per day. The ASGP-R may facilitate entry of the virus into the cell by providing a cell-surface receptor for glycoproteins on the virion. The glycosylation patterns of insect and mammalian cells differ, with the carbohydrate moieties on the surface of the virion produced in insect cells lacking terminal sialic acid. Those carbohydrate moieties may mediate internalization and trafficking of the virion. In addition to the ASGP-R, other galactose-binding lectins that exist in mammals (see, e.g., Jung et al., 1994, J. Biochem. (Tokyo) 116:547-553) may mediate uptake of the virus.

If desired, the cell to be infected can be modified to facilitate entry of the baculovirus into the cell. For example, ASGP-R can be expressed on the surface of a cell to be infected by the virus (e.g., baculovirus). The genes encoding the ASGP-R have been cloned (Spiess et al., 1985, J. Biol. Chem. 260:1979 and Spiess et al., 1985, Proc. Natl. Acad. Sci. 82:6465), and standard methods (e.g., retroviral, adeno-associated virus, or adenoviral vectors or chemical methods) can be used for expression of the ASGP-R in the cell to be infected by a virus. Other suitable mammalian lectins can be substituted for the ASGP-R in such methods (see, e.g., Ashwell et al., 1982, Ann. Rev. Biochem. 51:531-534). Other receptors for ligands on the virion, such as receptors for insect carbohydrates or the CD4 receptor for HIV, can also be expressed on the

surface of the mammalian cell to be infected to facilitate infection (see, e.g., Monsigny et al., 1979, Biol. Cellulaire 33:289-300).

Entry into the cell also can be facilitated by modifying the virion, e.g., through chemical means, to enable the virion to bind to other receptors on the mammalian cell (see, e.g., Neda, et al., 1991, J. Biol. Chem. 266:14143-14146 and Burns et al., 1993, Proc. Natl. Acad. Sci. 90:8033-8037). Alternatively, the glycosylation patterns and levels of baculovirus can be modified by growing the virus on Ea4 cells, which are derived from *Estigmena acrea* (e.g., as described by Rooney et al. in Nature Biotech). In addition, one can modify the virus such that it expresses mammalian glycosylation enzymes (Jarvis et al., 1996, Nature Biotech. 14:1288-1292).

II. Therapeutic Use of a Non-mammalian DNA Virus Expressing an Exogenous Gene

The discovery that a non-mammalian DNA virus efficiently expressed a lacZ reporter gene in several mammalian cells indicates that a non-mammalian DNA virus can be used therapeutically to express an exogenous gene in a cell of a mammal. For example, the method of the invention can facilitate expression of an exogenous gene in a cell of a patient for treatment of a disorder that is caused by a deficiency in gene expression. Numerous disorders are known to be caused by single gene defects (see Table 7), and many of the genes involved in gene deficiency disorders have been identified and cloned. Using standard cloning techniques (see, e.g., Ausubel et al., *Current Protocols in Molecular Biology*, John Wiley & Sons, (1989)), a non-mammalian virus can be engineered to express a desired exogenous gene in a mammalian cell (e.g., a human cell).

TABLE 7. EXAMPLES OF DISORDERS THAT CAN BE TREATED WITH THE INVENTION AND GENE PRODUCTS THAT CAN BE MANUFACTURED WITH THE INVENTION

	Gene Product	Disorder
	fumarylacetoacetate hydrolase	hereditary tyrosinemia
5	phenylalanine hydroxylase	phenylketonuria
	LDL receptor	familial hypercholesterolemia
	alpha-1 antitrypsin	alpha-1 antitrypsin deficiency
	glucose-6-phosphatase	glycogen storage diseases
	porphobilinogen deaminase	diseases caused by errors in porphyrin metabolism, e.g., acute intermittent porphyria
10	CPS-I, OTC, AS, ASL, or arginase	disorders of the urea cycle
	factors VIII & IX	hemophilia
	cystathione β -synthase	homocystinuria
	branched chain ketoacid decarboxylase	maple syrup urine disease
	albumin	hypoalbuminemia
15	isovaleryl-CoA dehydrogenase	isovaleric acidemia
	propionyl CoA carboxylase	propionic acidemia
	methyl malonyl CoA mutase	methylmalonyl acidemia
	glutaryl CoA dehydrogenase	glutaric acidemia
	insulin	insulin-dependent diabetes
20	β -glucosidase	Gaucher's disease
	pyruvate carboxylase	pyruvate carboxylase deficiency
	hepatic phosphorylase or phosphorylase kinase	glycogen storage diseases
	glycine decarboxylase, H-protein, or T-protein	non-ketotic hyperglycinemias
	Wilson's disease copper-transporting ATPase	Wilson's disease
25	Menkes disease copper-transporting ATPase	Menkes disease
	cystic fibrosis transmembrane conductance regulator	cystic fibrosis

The invention can also be used to facilitate the expression of a desired gene in a cell having no obvious deficiency. For example, the invention can be used to express insulin in a hepatocyte of a patient in order to supply the patient with insulin in the body. Other examples of proteins that can be

expressed in a mammalian cell (e.g., a liver cell) for delivery into the system circulation of the mammal include hormones, growth factors, and interferons. The invention can also be used to express a regulatory gene or a gene encoding a transcription factor (e.g., a VP16-tet repressor gene fusion) in a cell to control
5 the expression of another gene (e.g., genes that are operably-linked to a tet operator sequence; see, e.g., Gossen et al., 1992, Proc. Natl. Acad. Sci. 89:5547-5551). In addition, the invention can be used in a method of treating cancer by expressing in a cell a cancer therapeutic gene, such as a gene encoding a tumor suppressor (e.g., p53), tumor necrosis factor, thymidine
10 kinase, diphtheria toxin chimera, or cytosine deaminases (see, e.g., Vile and Russell, 1994, Gene Therapy 1:88-98).

Other useful gene products include RNA molecules for use in RNA decoy, antisense, or ribozyme-based methods of inhibiting gene expression (see, e.g., Yu et al., 1994, Gene Therapy 1:13-26). If desired, the invention can be
15 used to express a gene, such as cytosine deaminase, whose product will alter the activity of a drug or prodrug, such as 5-fluorocytosine, in a cell (see, e.g., Harris et al., 1994, Gene Therapy 1: 170-175). Methods such as the use of ribozymes, antisense RNAs, transdominant repressors, polymerase mutants, or core or surface antigen mutants can be used to suppress hepatitis viruses (e.g.,
20 hepatitis virus A, B, C, or D) in a cell. Other disorders such as familial hemochromatosis can also be treated with the invention by treatment with the normal version of the affected gene.

Preferred genes for expression include those genes that encode proteins that are expressed in normal mammalian cells (e.g., hepatocytes or lung cells).
25 For example, genes encoding enzymes involved in the urea cycle, such as the genes encoding carbamoyl phosphate synthetase (CPS-I), ornithine transcarbamylase (OTC), arginosuccinate synthetase (AS), arginosuccinate lyase (ASL), and arginase are useful in this method. All of these genes have been cloned (for OTC, see Horwich et al., 1984, Science 224:1068-1074 and Hata et al., 1988, J. Biochem (Tokyo) 103:302-308; for AS, see Bock et al., 1983, Nucl. Acids Res. 11:6505; Surh et al., 1988, Nucl. Acids Res. 16:9252; and
30 Dennis et al., 1989, Proc. Natl. Acad. Sci. 86:7947; for ASL, see O'Brien et

al., 1986, Proc. Natl. Acad. Sci. 83:7211; for CPS-I, see Adcock et al., 1984, (Abstract) Fed. Proc. 43:1726; for arginase, see Haraguchi et al., Proc. Natl. Acad. Sci. 84:412). Subcloning these genes into a baculovirus can be readily accomplished with common techniques.

5 The therapeutic effectiveness of expressing an exogenous gene in a cell can be assessed by monitoring the patient for known signs or symptoms of a disorder. For example, amelioration of OTC deficiency and CPS deficiency can be detected by monitoring plasma levels of ammonium or orotic acid. Similarly, plasma citrulline levels provide an indication of AS deficiency, and
10 ASL deficiency can be followed by monitoring plasma levels of arginosuccinate. Parameters for assessing treatment methods are known to those skilled in the art of medicine (see, e.g., Maestri et al., 1991, J. Pediatrics, 119:923-928).

 The non-mammalian DNA virus (e.g., baculovirus) can be formulated
15 into a pharmaceutical composition by admixture with a pharmaceutically acceptable non-toxic excipient or carrier (e.g., saline) for administration to a mammal. In practicing the invention, the virus can be prepared for use in parenteral administration (e.g., for intravenous injection (e.g., into the portal vein)), intra-arterial injection (e.g., into the femoral artery or hepatic artery),
20 intraperitoneal injection, intrathecal injection, or direct injection into a tissue or organ (e.g., intramuscular injection). In particular, the non-mammalian virus can be prepared in the form of liquid solutions or suspensions in conventional excipients. The virus can also be prepared for intranasal or intrabronchial administration, particularly in the form of nasal drops or aerosols in
25 conventional excipients. If desired, the virus can be sonicated in order to minimize clumping of the virus in preparing the virus.

 In practicing the invention, the virus can be used to infect a cell outside of the mammal to be treated (e.g., a cell in a donor mammal or a cell *in vitro*), and the infected cell then is administered to the mammal to be treated. In this
30 method, the cell can be autologous or heterologous to the mammal to be treated. For example, an autologous hepatocyte obtained in a liver biopsy can be used (see, e.g., Grossman et al., 1994, Nature Genetics 6:335). The cell can

then be administered to the patient by injection (e.g., into the portal vein). In such a method, a volume of hepatocytes totaling about 1% - 10% of the volume of the entire liver is preferred. Where the invention is used to express an exogenous gene in a liver cell, the liver cell can be delivered to the spleen, and the cell can subsequently migrate to the liver *in vivo* (see, e.g., Lu et al., 1995, Hepatology 21:7752-759). If desired, the virus may be delivered to a cell by employing conventional techniques for perfusing fluids into organs, cells, or tissues (including the use of infusion pumps and syringes). For perfusion, the virus is generally administered at a titer of 1×10^6 to 1×10^{10} pfu/ml (preferably 1×10^9 to 1×10^{10} pfu/ml) in a volume of 1 to 500 ml, over a time period of 1 minute to 6 hours. If desired, multiple doses of the virus can be administered to a patient intravenously for several days in order to increase the level of expression as desired.

The optimal amount of virus or number of infected cells to be administered to a mammal and the frequency of administration are dependent upon factors such as the sensitivity of methods for detecting expression of the exogenous gene, the strength of the promoter used, the severity of the disorder to be treated, and the target cell(s) of the virus. Generally, the virus is administered at a multiplicity of infection of about 0.1 to 1,000; preferably, the multiplicity of infection is about 5 to 100; more preferably, the multiplicity of infection is about 10 to 50.

III. Examples of Use of a Non-mammalian Virus to Express an Exogenous Gene *In Vivo*

The following examples demonstrate that a non-mammalian DNA virus can be used to express an exogenous gene in a cell *in vivo*. These examples also demonstrate that *in vivo* gene expression can be achieved by administering the virus by intravenous injection, intranasal administration, or direct injection of the virus into the targeted tissue. The first example demonstrates expression of an exogenous gene in brain cells *in vivo*. The second example provides evidence of expression of an exogenous gene in liver, following intravenous injection of the virus. In the third example, expression of the exogenous gene

is detected in skin after topical application of the Z4 virus to injured skin. In the remaining examples, a virus carrying an exogenous gene was injected directly into an organ. These examples demonstrate *in vivo* expression of an exogenous gene in skin, liver, spleen, kidney, stomach, skeletal muscle, uterus, and pancreas.

Injection Into Portal Vein: For the first example, 0.5 ml of Z4 virus ($\approx 1.4 \times 10^9$ pfu/ml) was injected (at a rate of ≈ 1 ml/min) into the portal vein of a single rat. At approximately 72 hours after infection, *lacZ* expression was detectable in at least one liver cell of the cryosections that were examined by conventional histochemical methods. The efficiency of expression may be increased by any one, or a combination of, the following procedures: (1) pre-treating the animal with growth factors; (2) partial hepatectomy, (3) administration of immunosuppressants to suppress any immune response to the virus; (4) use of a higher titer or dose of the virus; (5) infusion of the virus by surgical perfusion to the liver (e.g., in order to limit possible non-specific binding of the virus to red blood cells); and/or (6) sonication of the virus to minimize clumping of the virus.

Expression in Brain: For the second example, a 2 μ l sample of Z4 virus (at a titer of 4.8×10^{10} pfu/ml) was injected, using stereotactic procedure, into the olfactory bulb in the brain of an anesthetized adult rat. The virus was injected slowly (over a 30 minute time period) to avoid compressing the brain tissue. At 1 day post-injection, the rat was euthanized, and the brain tissue was processed for detection of expression of the exogenous *lacZ* gene by X-gal histochemistry. Injection of the Z4 virus into the brain resulted in *in vivo* expression of *lacZ*, as was evidenced by patches of cells that were strongly stained blue. More than 10^4 cells were stained blue upon injection of approximately 10^7 pfu. These data thus indicate that an exogenous gene can be expressed in the brain of a mammal by injecting into the brain a non-mammalian DNA virus whose genome includes the exogenous gene.

Topical Application and Expression in Skin: This example demonstrates that topical application of the Z4 virus to abraded skin of a mouse can result in expression of a heterologous gene in the skin. These experiments involved four

5 differently-treated areas on the skin of a mouse. Two of the areas (an abraded and a non-abraded area) were treated with phosphate-buffered saline. The other two areas (an abraded and a non-abraded area) were treated with the Z4 virus (50 μ l at 4.8×10^{10} pfu/ml). After treatment, each area of the skin was cut into sections using a cryostat.

10 Topical application of the Z4 virus (50 μ l at 4.8×10^{10} pfu/ml) to injured skin of a mouse resulted in expression of the exogenous gene in nearly 100% of the cells of the basal layer of the epidermis. Staining of deeper structures was not detected. In one cryostat section, various areas of the epidermis were stained in multiple sections. In a second cryostat section, occasional blue cells were present. In a third cryostat section, patches of staining were detected, and in a fourth cryostat section, the staining was nearly continuous and very dark. Although the pattern of gene expression varied slightly between the four cryostat sections obtained from this area of skin, the example demonstrates that
15 topical application of the Z4 virus to abraded skin consistently resulted in expression of the heterologous gene in skin.

Injection Into a Tissue or Organ: In the following examples, expression of an exogenous gene was detected *in vivo* after a non-mammalian DNA virus carrying the gene was injected directly into four distinct organs. For these
20 examples, the Z4 virus was prepared from 1 L of Z4-infected (moi of 0.5) Sf9 cells grown in spinner culture in serum-free medium. The cells and debris were removed by centrifuging the cell culture at 2000 rpm for 10 minutes. The virus was pelleted by centrifugation through a sucrose cushion in an SW28 rotor at 24,000 rpm for 75 minutes. For preparation of this virus stock, 33 ml
25 of cleared virus was layered over a 3 ml sucrose cushion (27% sucrose (w/v) in 10 mM Tris-HCl (pH 7.5), 1 mM EDTA (TE)). The virus was resuspended by overnight incubation at 4°C in 0.3 ml TE per tube. The virus was purified by banding in a 20-60% sucrose (w/v in TE) gradient in SW41 tubes that were centrifuged at 38,000 rpm for 75 minutes. The virus bands were collected with
30 a syringe and pelleted in SW50.1 rotor centrifuged at 30,000 rpm for 60 minutes. The virus pellet was resuspended in a total of 0.7 ml PBS by

overnight incubation at 4°C. The titer of the concentrated Z4 stock, as determined in a conventional plaque assay, was 4.8×10^{10} pfu/ml.

To assay for gene expression *in vivo*, the Z4 virus was administered Balb/c female mice by direct injection of a 50 μ l aliquot of the concentrated virus (2.4 x 10⁹ pfu total) into either the liver, spleen, kidney, muscle, uterus, pancreas, or skin of a mouse. Surgery was required for administration to liver, spleen and kidney. To spread the virus throughout an organ, the 50 μ l virus sample was injected into two or three sites in an organ. A 50 μ l sample of PBS was used as a negative control. For assaying gene expression in the liver, only one lobe of the liver was injected, and a separate mouse received the PBS injection as a negative control. For assaying gene expression in the spleen, an uninjected mouse served as a negative control. For assaying gene expression in kidney, muscle, and skin, contralateral controls were performed (the Z4 virus was injected into the right side of the organ, and PBS was injected into the left of the organ). For assaying expression in muscle, the virus was injected into the tibialis anterior hind leg muscle after shaving the mouse. For assaying expression in skin, the abdomen of the mouse was shaved, and 50 μ l of Z4 virus were injected into a marked section of the abdomen. At 24 hours post-injection, the mice were sacrificed and dissected. The Z4- and PBS-injected organs were frozen in liquid nitrogen, and 7 μ m thin sections were prepared using a cryostat (Reichert-Jung Cryocut 1800). β -galactosidase activity was measured by fixing the thin sections and staining with X-gal, as described above. Each of the organs that received the Z4 virus expressed the exogenous *lacZ* gene *in vivo*. In each case, the PBS negative control did not promote expression of the exogenous gene.

Injection and Expression in Skin: In this example, *in vivo* expression of the exogenous *lacZ* gene of Z4 was observed in mouse skin after injection of 2.4 x 10⁹ pfu into the skin. A high level of expression (over 25% of cells within the area of injection) was achieved in the dermis after subcutaneous injection of the virus. Although the muscle layer was predominantly unstained, positive staining of some skeletal muscle fibers was observed. As a negative control, PBS was injected into the skin. Although some staining was observed

in the sebaceous glands, it is most probably due to the presence of bacteria. A low level of staining was also detected in the dermis. Similar results were obtained when the Z4 virus was applied topically to uninjured (non-abraded) skin, although no clear epidermal staining was detected. Nonetheless, these data indicate that the Z4 virus can be used to express a heterologous gene in the skin of a mammal when the virus is injected subcutaneously into the mammal.

Expression in Liver: In this example, expression of the exogenous gene was detected in liver. Blue coloration, indicative of β -galactosidase expression, was detected in multiple areas of the injected lobe. Although the most intense coloration was at the point of injection, the internal areas of the liver sections exhibited the blue coloration that is indicative of gene expression. Expression of the exogenous gene appeared to be detected both in hepatocytes and Kupffer cells of the lobes that received the Z4 virus. In contrast, uninjected lobes from the same liver were negative. These results thus indicate that an exogenous gene can be expressed in a liver cell by injecting into the liver a non-mammalian DNA virus encoding the gene.

Expression in Spleen: In this example, thin sections of the spleen were assayed for gene expression following injection of the virus carrying the exogenous gene into the spleen. Spleen cells that had received the Z4 virus *in vivo* expressed the *lacZ* gene. The blue coloration was detected in cells located throughout the entire spleen. The intensity of blue coloration obtained with spleen cells was less than the intensity obtained with liver cells. Nonetheless, the blue coloration was indicative of significant expression of the exogenous gene. No blue coloration was detected in a spleen that did not receive the virus. These data thus indicate that an exogenous gene can be expressed in a spleen cell *in vivo* upon injection of a non-mammalian DNA virus whose genome carries the gene.

Expression in Kidney: In this example, *in vivo* expression of an exogenous gene was detected in a kidney that was injected with Z4 as described above. The Z4-injected kidney displayed clear blue coloring that is indicative of *lacZ* expression; in contrast, a PBS-injected control kidney displayed no blue

coloration. The blue coloration was primarily around the edges of the sections of the kidney. Indirect immunofluorescence also indicated that the viral particles were concentrated in the edges of the sections, providing a correlation between gene expression and localization of the virus. These data thus indicate that a non-mammalian DNA virus can be used to express an exogenous gene in a kidney cell *in vivo*.

Expression in Stomach: In this example, the Z4 virus (50 μ l) was injected into the center of the stomach of Balb/C mice. The animals were sacrificed on the day following injection, and the stomachs were frozen in liquid nitrogen, and cryostat sectioned and stained as previously described. Cell transfection was observed in gastric mucosal and muscle cells. Positive staining was detected in glands, with most staining occurring at the bases of the glands. These observations indicate that a non-mammalian DNA virus can be used to express a heterologous gene in the stomach of mice. In these experiments, blue staining was also detected in the lumen. The blue coloration in that particular region may result from bacteria in the gut, rather than expression from the virus.

Expression in Skeletal Muscle: In this example, *in vivo* expression of the exogenous *lacZ* gene of Z4 was detected in muscle after direct injection of virus into the tibialis anterior. Blue coloration was found only in discrete loci in the muscle, and the coloration was not as intense or widespread as the coloration observed in liver, spleen, or skin. Nonetheless, the blue coloration was significant, indicating that a non-mammalian DNA virus can be used to express an exogenous gene in muscle *in vivo*.

Expression in Uterus: In this example, expression of the *lacZ* reporter gene was detected in the uterus. A 50 μ l aliquot of the Z4 virus (2.4×10^9 pfu) was injected directly into the uterus of a mouse. The animal was sacrificed on the day following injection, and cryostat sections were prepared as previously described. Staining of the sections with X-gal produced blue coloration in an area of the uterus with little tissue disruption. The positive cells were mostly endometrial stromal cells, rather than gland elements. These

data indicate that a non-mammalian DNA virus can be used to express a heterologous gene in the uterus of a mammal.

Expression in Pancreas: This example demonstrates that a non-mammalian DNA virus can be used to express a heterologous gene in the pancreas of a mammal. A 50 μ l aliquot of the Z4 virus (2.4×10^9 pfu) was injected directly into the pancreas of a mouse. On the day following injection, the mouse was sacrificed, and the pancreas was stained with X-gal according to conventional methods. Large areas of positive cells were detected, indicating that the Z4 virus successfully expressed the *lacZ* gene in the pancreas.

Summary: In sum, these examples demonstrate that a non-mammalian DNA virus (e.g., a baculovirus) can be used to express an exogenous gene in a mammalian cell *in vivo*. These examples employed several distinct animal model systems and methods of administering the virus. In each and every case, the non-mammalian DNA virus successfully expressed the exogenous gene *in vivo*. These data thus provide support for the assertion that a non-mammalian DNA virus can be used to express an exogenous gene in other, non-exemplified cells *in vivo*. In addition, in at least some tissues, the level of expression *in vivo* was, surprisingly, higher than the level that would have been predicted from the corresponding *in vitro* experiments (e.g., the brain versus cultured neurons). All of these examples provide evidence of the *in vivo* utility of the invention.

PART B: AN ALTERED COAT PROTEIN ENHANCES THE ABILITY OF A NON-MAMMALIAN DNA VIRUS TO EXPRESS AN EXOGENOUS GENE IN A MAMMALIAN CELL

Now that it has been demonstrated that a non-mammalian DNA virus can be used to express an exogenous gene in a mammalian cell, the following examples are provided to demonstrate that an altered coat protein enhances the ability of the non-mammalian DNA virus to express the exogenous gene in a mammalian cell.

Construction of Baculovirus Transfer Plasmid pVGZ3

These examples employ a baculovirus that has been engineered to express a vesicular stomatitis virus glycoprotein G (VSV-G) as an altered coat

protein. The baculovirus transfer plasmid pVGZ3 is a derivative of the baculovirus transfer plasmid Z4, which was used in many of the examples summarized above.

5 The baculovirus transfer plasmid pVGZ3 was constructed by inserting expression cassettes encoding VSV-G and the reporter gene *lacZ* into the baculovirus transfer vector BacPAK9 (Clontech; Palo Alto, CA). To produce this transfer plasmid, cDNA encoding VSV-G was excised as a 1,665 bp *Bam*HI fragment from pLGRNL (Burns, 1993, Proc. Natl. Acad. Sci. 90:8033-8037). The resulting plasmid was termed VSVG/BP9 (Fig. 18). The RSV
10 LTR, *lacZ* gene, and SV40 polyadenylation signal were excised from the Z4 transfer plasmid using *Bgl*II and *Bam*HI to produce a 4,672 bp fragment. This fragment was inserted into the *Bgl*II site of VSVG/BP9 such that the *lacZ* gene was positioned downstream from the VSV-G gene to produce the VGZ3 transfer plasmid (Fig. 19). In VGZ3, the directions of transcription of the
15 VSV-G and *lacZ* genes are convergent. In other words, the promoters lie at opposite ends of the inserted sequences, with the *lacZ* and VSV-G genes being transcribed towards each other. The SV40 polyA site is bidirectional and used by both the VSV-G and *lacZ* genes.

Because the VSV-G gene in this transfer plasmid is operably linked to a
20 baculovirus polyhedrin promoter, the plasmid offers the advantage of high levels of expression of the VSV-G gene in insect cells and relatively low levels of expression in mammalian cells. This pattern of expression of the altered coat protein is desirable because the altered coat protein is produced efficiently in insect cells, where the non-mammalian DNA virus having an protein is
25 manufactured before the virus is delivered to a mammalian cell. Producing the virus in insect cells before using the virus to infect mammalian cells obviates concerns about expressing a viral coat protein (e.g., VSV-G) in mammalian cells.

A schematic representation of a budding virus having an altered coat
30 protein is provided in Fig. 20. Once the virus infects a mammalian cell, expression of the altered coat protein is not desirable. Therefore, the use of an non-mammalian (e.g., insect) virus promoter, which is not active in mammalian

cells, is preferable for driving expression of the altered coat protein. By contrast, the exogenous gene of interest (here, the *lacZ* reporter gene) is operably linked to an RSV LTR. As is desired, expression of a gene driven by the RSV LTR is obtained both in insect cells and in mammalian cells.

5 Standard procedures were used to produce the recombinant baculovirus VGZ3 from the pVGZ3 transfer plasmid, described above. Briefly, Sf9 cells were co-transfected, according to the manufacturer's instructions, by lipofection with pVGZ3 and the baculovirus genomic DNA BacPAK6 (Clontech; Palo Alto, CA) that had been digested with *Bsu36I*. The virus was plaque purified
10 and amplified according to standard techniques. Using a conventional plaque assay on Sf9 cells, the viral titer was determined to be 3.1×10^7 pfu/ml.

Enhanced Expression in HepG2, Vero, and HeLa Cells: This example demonstrates that VGZ3, the baculovirus having an altered coat protein, has an enhanced ability, relative to the Z4 virus described above, to express an
15 exogenous gene in a mammalian cell. To provide sensitivity in the assays, these experiments were performed under conditions in which the Z4 virus expresses an exogenous gene at relatively low levels. Three different cell types were used, including HepG2 cells, Vero cells (a kidney cell line), and HeLa
20 cells (a cervical carcinoma cell line). In these experiments, 1×10^5 cells of each cell type were, independently, seeded into multiple wells of 12-well culture plates. On the following day, the tissue culture medium was replaced with fresh medium, and the Z4 and VGZ3 viruses were added, separately, to the cells. The viruses were used at multiplicities of infection of 0, 1.25, 10,
25 and 80 (assuming that the cell number had doubled overnight), and each experiment was performed in duplicate. On the day following the addition of virus, the cells were harvested, and expression of the exogenous *lacZ* gene was detected by X-gal staining or by using a quantitative chemiluminescent β -galactosidase assay (Clontech; Palo Alto, CA). The results of these assays are presented in Table 8.

TABLE 8. USE OF A NON-MAMMALIAN DNA VIRUS HAVING AN ALTERED COAT PROTEIN TO ACHIEVE ENHANCED EXPRESSION OF AN EXOGENOUS GENE IN HEPG2 AND HELA CELLS

Cell Line	Virus	moi	Chemiluminescence Units	VGZ3 Superiority ^a
HepG2	Z4	80	16.4	
HepG2	VGZ3	80	180.2	11.0-fold
HeLa	Z4	80	0.02 ^b	
HeLa	VGZ3	80	1.75	>87.5-fold
HeLa	VGZ3	1.25	0.07	>224 fold ^c

^a Superiority was calculated as VGZ3 transduction units ÷ Z4 transduction units for each cell type.

^b 0.02 was the background level in the chemiluminescent assay.

^c The difference in moi (1.25 for VGZ3 and 80 for Z4) was accounted for in determining the VGZ3 superiority.

15 This example demonstrates that a baculovirus that is engineered to express a VSV glycoprotein G has an enhanced ability, relative to a baculovirus that lacks the altered coat protein, to express an exogenous gene in a mammalian cell. In this example, expression of the exogenous *lacZ* gene was detected in 100% of the HepG2 cells that were contacted with VGZ3 at an moi of 80. In contrast, under these conditions, expression of the exogenous gene was detected in only 15% of the cells that were contacted with the Z4 virus (a virus that does not have an altered coat protein). Accordingly, these data show that altering a coat protein enhances the ability of a virus to express an exogenous gene in a mammalian cell.

25 Enhanced expression of the exogenous gene was also achieved with Vero cells: over 50% of the VGZ3-treated cells turned blue, whereas only 5-10% of the Z4-treated cells turned blue upon staining with X-gal. Further evidence that the altered coat protein enhances the ability of a virus to express an exogenous gene in a mammalian cell comes from a relatively sensitive assay employing HeLa cells. Under the conditions employed in this example, the Z4 virus does

not efficiently express an exogenous gene in HeLa cells. No blue cells were detected at an moi of 80, indicating that the frequency of gene expression was less than 1×10^5 . By contrast, approximately 3% of the HeLa cells treated with the VGZ3 virus were blue, indicating that the efficiency of gene
5 expression was 3×10^{-2} , which is 12,000-fold better than the efficiency obtained with Z4. With the VGZ3 virus, blue HeLa cells were also detected at the low moi of 1.25, whereas no blue cells were detected at the moi with the Z4 virus.

In sum, these data indicate that an altered coat protein on a non-
10 mammalian DNA virus can enhance the ability of that virus to infect and express a gene in a mammalian cell. In addition, employment of an altered coat protein allows the virus to infect a cell at a lower moi. Thus certain cells that appear to be refractive to the virus at a given moi (e.g., HeLa cells) can be infected with a virus having an altered coat protein, thereby expanding the
15 apparent host range of the virus. A virus having an altered coat protein thus offers the advantage of permitting expression of an exogenous gene at a low moi, relative to the moi needed with a virus that lacks an altered coat protein.

As was described above for the Z4 virus, exogenous gene expression in cells treated with the VGZ3 virus results from *de novo* gene expression within
20 the mammalian cell. Both cycloheximide, which inhibits protein synthesis, and chloroquine, which inhibits endosome acidification, separately inhibited β -galactosidase expression in VGZ3-treated mammalian cells (data not shown). Thus, β -galactosidase detected in the mammalian cells in these experiments can be attributed to *de novo* protein synthesis within the mammalian cell.

25 Enhanced Expression in PC12 Cells: In this example, the VGZ3 virus is shown to increase the level of exogenous gene expression in rat cortical neuronal cells, relative to the level obtained with the Z4 virus. Using conventional methods, PC12 cells were plated at 10,000 cells/well in a 24-well dish. On day 1, the cell culture medium (DMEM containing 5% FBS and 10%
30 horse serum) was replaced with fresh cell culture medium that also contained 50 ng/ml of nerve growth factor (NGF). Fresh NGF-containing-medium was again added at days 3 and 5. On day 6, the cells were infected with various

dilutions of Z4 virus and VGZ3 virus, as shown in Table 9. At day 7, the cells were fixed and stained for β -galactosidase by immunocytochemistry using an anti- β -galactosidase antibody (available from 5'->3' Inc.). Under these conditions, fewer than 1% of the PC12 cells infected with 2×10^8 pfu of Z4 expressed the exogenous gene, and approximately 17.5% of the cells infected with 1×10^7 pfu of VGZ3 expressed the exogenous gene. Accordingly, these data provide additional evidence that a virus having an altered coat protein has a superior ability to express an exogenous gene in a mammalian cell.

Enhanced Expression in Primary Rat Cortical Cells

This example demonstrates that a virus having an altered coat protein provides enhanced expression of an exogenous gene in a primary cultures of rat cortical cells, as compared with a virus lacking the altered coat protein. For this example, cultures of rat cortical cells were prepared and infected as described above under "Expression in Cortex Cultures." The Z4 and VGZ3 viruses were used at the moi shown in Table 9 (note: the moi of Z4 was approximately 10-fold higher than the moi of VGZ3). When the number of blue cells obtained is compared with the moi of Z4 or VGZ3 used, it becomes apparent that the VGZ3 virus is more efficient at expressing the exogenous gene in the mammalian cells than is the Z4 virus. In addition, this example demonstrates that a non-mammalian DNA virus having an altered coat protein can direct exogenous gene expression in primary rat neurons (in addition to the cell line PC12, as shown above).

TABLE 9. USE OF A NON-MAMMALIAN DNA VIRUS HAVING AN ALTERED COAT PROTEIN TO ACHIEVE ENHANCED EXPRESSION OF AN EXOGENOUS GENE IN PRIMARY CULTURES OF RAT CORTICAL CELLS

Virus	1 μ l	2 μ l	5 μ l	10 μ l	50 μ l	100 μ l
Z4	moi=1 ^a no blue cells	moi=2 no blue cells	moi=5 ≈5 blue cells	moi=10 ≈20 blue cells	moi=50 ≈500 blue cells	moi=100 ≈2200 blue cells ≈0.75%
VGZ3	moi=0.1 no blue cells	moi=0.2 no blue cells	moi=0.5 no blue cells	moi=1 ≈10 blue cells	moi=5 ≈200 blue cells	moi=10 ≈60 blue cells ^b
PBS				no blue cells	no blue cells	no blue cells

^a The moi's were estimated based on the number of cells plated the day before infection.

^b Because of cell death occurring in this well, fewer stained cells were detected. Nonetheless, the percentage of blue cells was high.

Enhanced Expression in HepG2, HuH7, HeLa, WISH, A549, VERO, CHO, and Balb/c 3T3 Cells: Further data showing that an altered coat protein enhances the ability of a non-mammalian DNA virus to direct expression of an exogenous gene in mammalian cells is provided by this example. Here, a variety of cells were infected with the VGZ3 baculovirus. The methods employed in these experiments first are described.

Cells: The human hepatoma lines HepG2 and HuH7, the human cervical carcinoma line HeLa, the human amniotic cell line WISH, the human lung carcinoma A549, the African green monkey kidney line VERO, the hamster epithelial line CHO and the mouse embryonic fibroblast line Balb/c 3T3 were all obtained from ATCC. All mammalian cells were grown in Dulbecco's Modified Eagle's Medium (GibcoBRL, Grand Island, NY) with 10% fetal bovine serum and 4 mM glutamine (BioWhittaker, Walkersville, MD), except for WISH cells, which were grown in MEM with Hanks's salts (GibcoBRL), 20% fetal bovine serum and 4 mM glutamine.

Infection and Reporter Gene Assay: Cells were seeded at 2×10^5 cells per well in 12-well plates. After the cells attached to the plastic, the cells were rinsed with medium and fresh complete medium was added. Viral infection was performed by adding virus to the medium at the indicated multiplicities of infection (moi). Following an 18-24 hour incubation at 37°C in 5% CO₂, cells were stained with X-gal to visualize β -galactosidase-expressing cells or cell lysates were taken and β -galactosidase activity quantitated by a luminescent β -galactosidase assay (Clontech catalog # K2048-1) according to the manufacturer's instructions.

Results: The use of the VGZ3 virus enhances exogenous gene expression, as compared with the level of gene expression obtained with the Z4 virus. X-gal staining of infected cells in culture indicated that an approximately 10-fold higher percentage of HepG2 cells expressed the exogenous gene following infection with VGZ3, as compared with the Z4 virus (data not shown). In addition, the intensity of the blue staining has greater in the VGZ3-treated cells, suggesting that a higher level of gene expression within the VGZ3-infected cells. Enhanced gene expression was also detected when the VGZ3 virus was used to infect HeLa cells. At an moi of 100, the Z4 virus produced few blue cells per well (approximately 1-5 cells), while approximately 10% of the VGZ3 cells stained blue with X-gal.

The results of a quantitative assay of β -galactosidase expression are shown in Fig. 22. At each moi tested, the level of β -galactosidase expression in HepG2 cells treated with VGZ3 was roughly 10-fold higher than the level obtained with the Z4 virus. The difference in transduction efficiency between the Z4 virus and the VGZ3 virus was even more notable in HeLa cells. At an moi of 1 or 10, no β -galactosidase activity above the background levels was detected with the Z4 virus. In contrast, β -galactosidase activity was detectable in HeLa cells treated with VGZ3 at an moi of 1. When the Z4 virus was used at an moi of 100, β -galactosidase activity just above background levels was detected.

When the VGZ3 virus was used at an moi of 100, the level of β -galactosidase activity detected in HeLa cells was approximately 350 times greater than the level detected in Z4-treated cells.

A panel of 8 different cell lines was used to compare the transduction efficiencies of the Z4 and VGZ3 viruses at an moi of 50. At this low moi, exogenous gene expression is not detected in certain of the cell lines treated with the Z4 virus, as shown in Table 10. In contrast, the VGZ3 virus led to detectable levels of exogenous gene expression in all of the cell lines at an moi of 50. In sum, these data provide further evidence that an altered coat protein enhances exogenous gene expression from a non-mammalian DNA virus.

TABLE 10: B-GALACTOSIDASE ACTIVITY IN Z4- AND VGZ3-TREATED CELLS

β -galactosidase activity ^a		
Cells	Z4-treated ^b	VGZ3-treated
HepG2	6.62	58.21
HuH7	4.46	42.49
HeLa	0.05	2.67
WISH	0.00	1.85
A549	0.22	46.34
VERO	0.58	6.38
CHO	0.00	2.33
3T3	0.02	2.01

^a For each cell line, the β -galactosidase activity in uninfected cells was determined, and this value was subtracted from the raw numbers for β -galactosidase activity in Z4-treated and VGZ3-treated cells. Each data point represents the average of three samples.

^b All Z4 and VGZ3 treatments were at an moi of 50.

Other Embodiments

Non-mammalian viruses other than the above-described *Autographa californica* viruses can be used in the invention; such viruses are listed in Table 1. Nuclear polyhedrosis viruses, such as multiple nucleocapsid viruses (MNPV) or single nucleocapsid viruses (SNPV), are preferred. In particular, *Choristoneura fumiferana* MNPV,

Mamestra brassicae MNPV, *Buzura suppressaria* nuclear polyhedrosis virus, *Orygia pseudotsugata* MNPV, *Bombyx mori* SNPV, *Heliothis zea* SNPV, and *Trichoplusia ni* SNPV can be used.

Granulosis viruses (GV), such as the following viruses, are also included among those that can be used in the invention: *Cryptophlebia leucotreta* GV, *Plodia interpunctella* GV, *Trichoplusia ni* GV, *Pieris brassicae* GV, *Artogeia rapae* GV, and *Cydia pomonella* granulosis virus (CpGV). Also, non-occluded baculoviruses (NOB), such as *Heliothis zea* NOB and *Oryctes rhinoceros* virus can be used.

Other insect (e.g., lepidopteran) and crustacean viruses can also be used in the invention. Further examples of useful viruses include those that have infect fungi (e.g., *Strongwellsea magna*) and spiders. Viruses that are similar to baculoviruses have been isolated from mites, Crustacea (e.g., *Carcinus maenas*, *Callinectes sapidus*, the Yellow Head Baculovirus of penaeid shrimp, and *Penaeus monodon*-type baculovirus), and Coleoptera. Also useful in the invention is the *Lymantria dispar* baculovirus.

If desired, the virus can be engineered to facilitate targeting of the virus to certain cell types. For example, ligands that bind to cell surface receptors other than the ASGP-R can be expressed on the surface of the virion. Alternatively, the virus can be chemically modified to target the virus to a particular receptor.

If desired, the cell to be infected can first be stimulated to be mitotically active. In culture, agents such as chloroform can be used to this effect; *in vivo*, stimulation of liver cell division, for example, can be induced by partial hepatectomy (see, e.g., Wilson, et al., 1992, J. Biol. Chem. 267:11283-11489). Optionally, the virus genome can be engineered to carry a herpes simplex virus thymidine kinase gene; this would allow cells harboring the virus genome to be killed by ganciclovir. If desired, the virus could be engineered such that it is defective in growing on its natural non-mammalian host cell (e.g., insect cell). Such strains of viruses could provide added safety and be propagated on a complementing packaging line. For example, a defective baculovirus could be made in which an immediate

early gene, such as IE1, has been deleted. This deletion can be made by targeted recombination in yeast or *E. coli*, and the defective virus can be replicated in insect cells in which the IE1 gene product is supplied in *trans*. If desired, the virus can be treated with neuraminidase to reveal additional

5 terminal galactose residues prior to infection (see, e.g., Morell et al., 1971, J. Biol. Chem. 246:1461-1467).

What is claimed is:

1. A method of expressing an exogenous gene in a mammalian cell, said method comprising:

a) introducing into the cell a non-mammalian DNA virus, wherein:

5 the virus has an altered coat protein, and the genome of the virus comprises an exogenous gene under the control of a promoter that induces expression of the exogenous gene in the cell; and

b) maintaining said cell under conditions such that said exogenous gene is expressed.

10 2. The method of claim 1, wherein said invertebrate virus is an insect virus.

3. The method of claim 2, wherein said insect virus is a baculovirus.

4. The method of claim 3, wherein said baculovirus is an *Autographa californica* multiple nuclear polyhedrosis virus.\

15 5. A method of treating a disorder in a mammal, comprising:

a) introducing into a cell a therapeutically effective amount of a non-mammalian DNA virus to produce an infected cell, wherein the virus has an altered coat protein, and the genome of the virus comprises an exogenous gene; and

20 b) maintaining said infected cell under conditions such that said exogenous gene is expressed in said mammal.

6. The method of claim 5, wherein the altered coat protein comprises a coat protein of a mammalian virus.

25 7. The method of claim 5, wherein the altered coat protein is a fusion protein.

8. The method of claim 5, wherein said virus is an insect virus.

9. The method of claim 8, wherein said insect virus is a baculovirus.

10. A method for treating a cancer in a mammal, said method comprising:

30 a) introducing into a cancerous cell of said mammal a non-mammalian DNA virus having an altered coat protein to produce an infected cell, wherein

the genome of said virus comprises a cancer-therapeutic gene encoding a protein selected from the group consisting of tumor necrosis factor, p53, thymidine kinase, diphtheria toxin chimeras, and cytosine deaminase; and

- 5 b) maintaining said infected cell in said mammal under conditions such that said cancer-therapeutic gene is expressed.

11. The method of claim 10, wherein the altered coat protein comprises a coat protein of a mammalian virus.

12. The method of claim 10, wherein the altered coat protein comprises a fusion protein.

- 10 13. The method of claim 10, wherein said cancerous cell is selected from the group consisting of hepatocytes, pancreas cells, lung cells, thyroid cells, thymus cells, prostate tissue cells, breast tissue cells, brain cells, neuronal cells, glial cells, skin cells, spleen cells, muscle cells, kidney cells, and bladder cells.

14. The method of claim 10, wherein the non-mammalian DNA virus is
15 an insect virus.

15. The method of claim 14, wherein said insect virus is a baculovirus.

16. A method for treating a neurological disorder in a mammal, said method comprising:

- 20 a) introducing into a cell a therapeutically effective amount of a non-mammalian DNA virus having an altered coat protein to produce an infected cell, wherein the genome of said virus comprises an exogenous gene encoding a therapeutic protein selected from the group consisting of nerve growth factor, hypoxanthine guanine phosphoribosyl transferase, tyrosine hydroxylase, dopadecarboxylase, brain-derived neurotrophic factor, and basic fibroblast
25 growth factor; and

 b) maintaining said infected cell under conditions such that said exogenous gene is expressed in said mammal.

17. The method of claim 16, wherein said virus is an insect virus.

18. The method of claim 17, wherein said insect virus is a baculovirus.

- 30 19. The method of claim 16, wherein said altered coat protein comprises a coat protein of a mammalian virus.

20. The method of claim 16, wherein the altered coat protein comprises a fusion protein.

21. A nucleic acid comprising:

a genome of a non-mammalian DNA virus;

5 an exogenous mammalian gene;

an exogenous mammalian-active promoter operably linked to said exogenous mammalian gene; and

a gene encoding an altered coat protein.

22. The nucleic acid of claim 21, wherein the gene encoding an altered coat protein encodes a coat protein of a mammalian virus.

23. The nucleic acid of claim 21, wherein the gene encoding the altered coat protein encodes a fusion protein.

24. The nucleic acid of claim 21, wherein said virus is an insect virus.

25. The nucleic acid of claim 21, wherein said insect virus is a baculovirus.

26. A nucleic acid comprising:

a genome of a non-mammalian DNA virus;

a gene encoding an altered coat protein;

20 an exogenous antisense RNA gene, the RNA transcribed from said exogenous antisense RNA gene being complementary to a nucleic acid of a gene that is expressed in a cell at an undesirably high level; and

an exogenous mammalian-active promoter, wherein said exogenous antisense RNA gene is operably linked to said promoter.

27. The nucleic acid of claim 26, wherein the gene encoding an altered coat protein encodes a coat protein of a mammalian virus.

28. The nucleic acid of claim 26, wherein the gene encoding the altered coat protein encodes a fusion protein.

29. The nucleic acid of claim 26, wherein said virus is an insect virus.

30. The nucleic acid of claim 29, wherein said insect virus is a baculovirus.

31. A cell that contains a nucleic acid, wherein said nucleic acid comprises:

a genome of a non-mammalian DNA virus;
a gene encoding an altered coat protein; and
an exogenous mammalian gene operably linked to an exogenous
mammalian-active promoter.

5 32. The cell of claim 31, wherein the gene encoding an altered coat
protein encodes a coat protein of a mammalian virus.

33. The cell of claim 31, wherein the gene encoding the altered coat
protein encodes a fusion protein.

34. The cell of claim 31, wherein said virus is an insect virus.

10 35. The cell of claim 34, wherein said insect virus is a baculovirus.

36. A nucleic acid comprising:

a genome of a non-mammalian DNA virus;

a gene encoding an altered coat protein;

15 an exogenous cancer therapeutic gene selected from the group
consisting of tumor necrosis factor genes, thymidine kinase genes, chimeric
diphtheria toxin genes, and cytosine diaminase genes; and

an exogenous mammalian-active promoter, wherein said exogenous
cancer therapeutic gene is operably linked to said promoter.

20 37. The nucleic acid of claim 36, wherein the gene encoding an altered
coat protein encodes a coat protein of a mammalian virus.

38. The nucleic acid of claim 36, wherein the gene encoding the altered
coat protein encodes a fusion protein.

39. The nucleic acid of claim 36, wherein said virus is an insect virus.

25 40. The nucleic acid of claim 39, wherein said insect virus is a
baculovirus.

41. A nucleic acid comprising:

a genome of a non-mammalian DNA virus

a gene encoding an altered coat protein;

30 an exogenous gene selected from the group consisting of RNA decoy
genes and ribozyme genes; and

an exogenous mammalian-active promoter operably linked to said
exogenous gene.

42. The nucleic acid of claim 41, wherein the gene encoding an altered coat protein encodes a coat protein of a mammalian virus.

43. The nucleic acid of claim 41, wherein the gene encoding the altered coat protein encodes a fusion protein.

5 44. The nucleic acid of claim 41, wherein the virus is an insect virus.

45. The nucleic acid of claim 44, wherein said insect virus is a baculovirus.

46. A pharmaceutical composition comprising:

(A) a pharmaceutically acceptable excipient and

10 (B) a non-mammalian DNA virus, wherein the genome of the virus comprises:

an exogenous mammalian gene;

an exogenous mammalian-active promoter operably linked to said exogenous mammalian gene; and

15 a gene encoding an altered coat protein.

47. The pharmaceutical composition of claim 46, wherein said virus is an insect virus.

48. The pharmaceutical composition of claim 47, wherein said insect virus is a baculovirus.

20 49. The pharmaceutical composition of claim 46, wherein the altered coat protein comprises the coat protein of a mammalian virus.

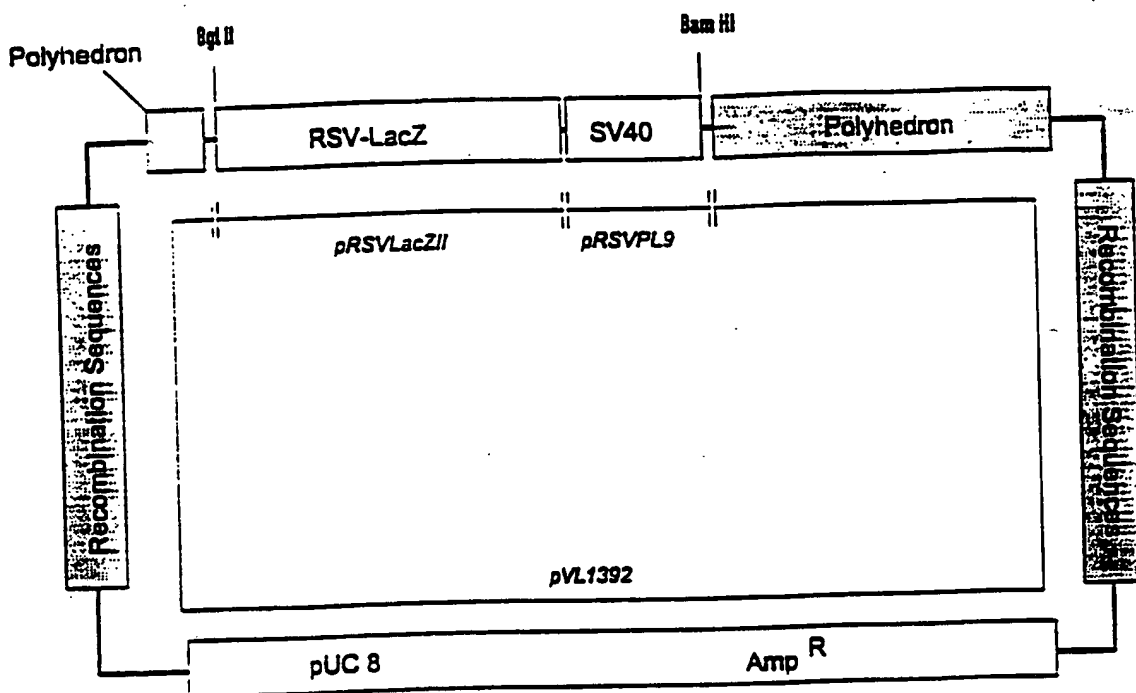
50. The pharmaceutical composition of claim 46, wherein the gene encoding the altered coat protein encodes a fusion protein.

25 51. A non-mammalian DNA virus having on its surface an altered coat protein, wherein the genome of the virus comprises an exogenous mammalian gene and an exogenous mammalian-active promoter operably linked to the exogenous mammalian gene.

52. The non-mammalian DNA virus of claim 51, wherein the genome of the virus further comprises a gene encoding an altered coat protein.

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FIG. 1
AcMNPV Transfer Plasmid Z4



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FIG. 2

AcMNPV Transfer Plasmid Z5

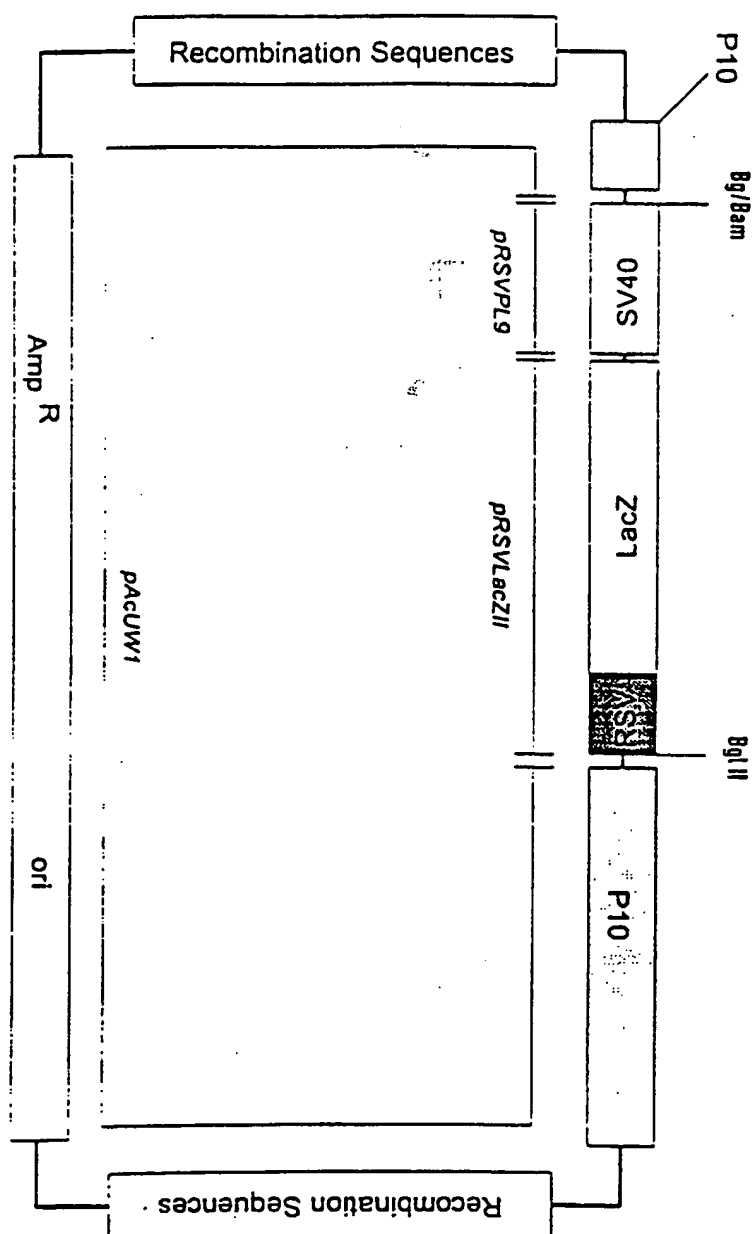
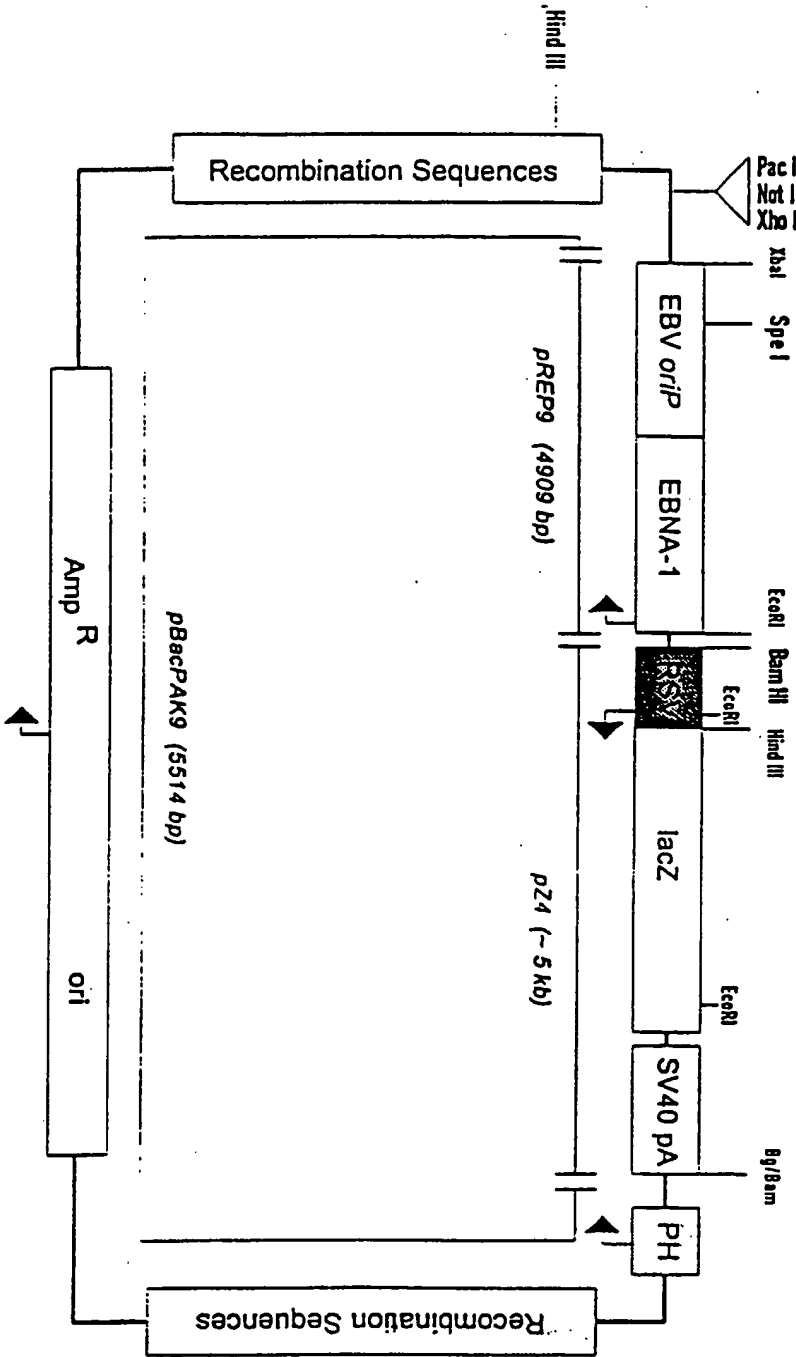


FIG. 3

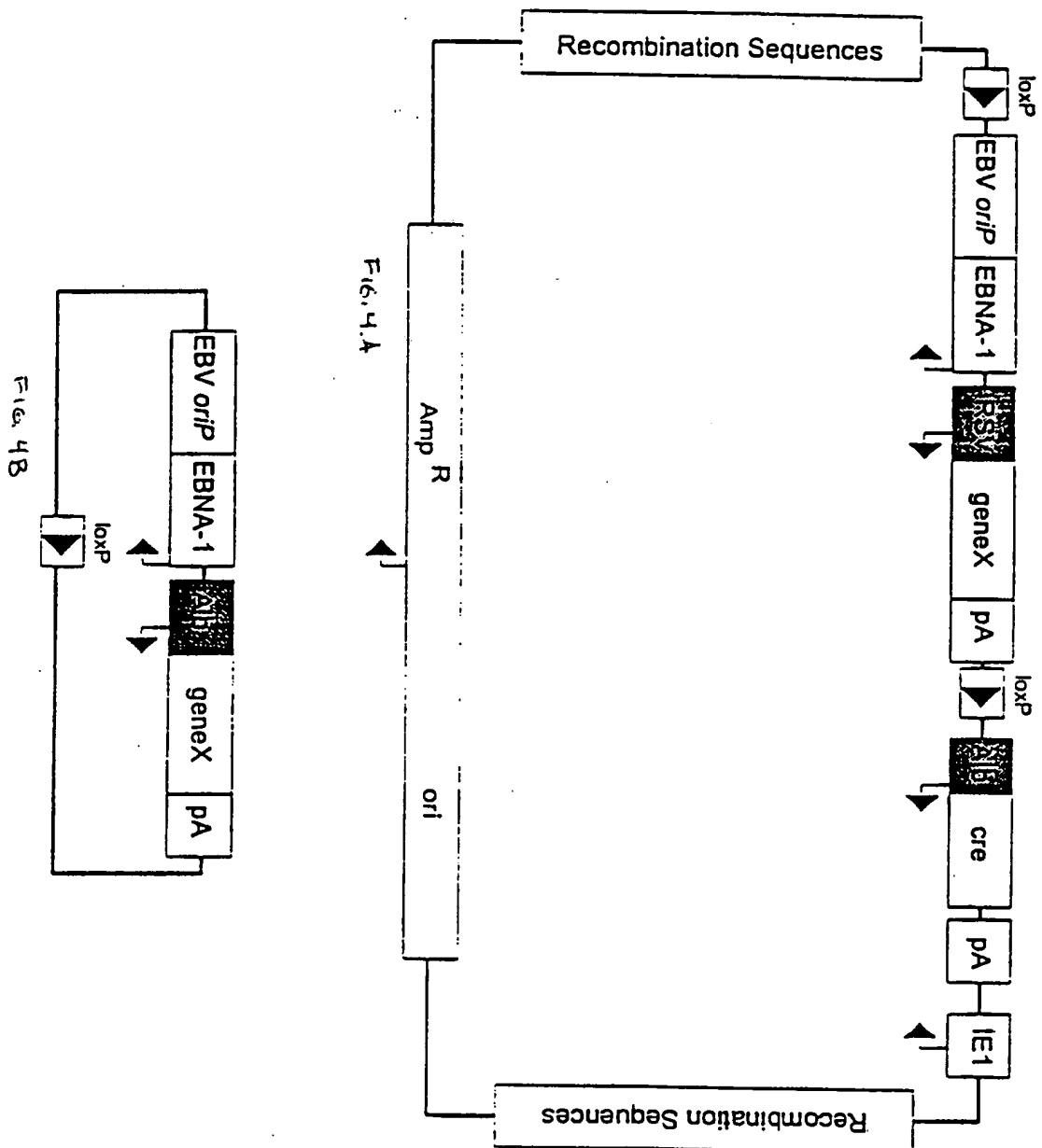
AcMNPV Transfer Plasmid Z-EBV #1



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FIG. 4A - B

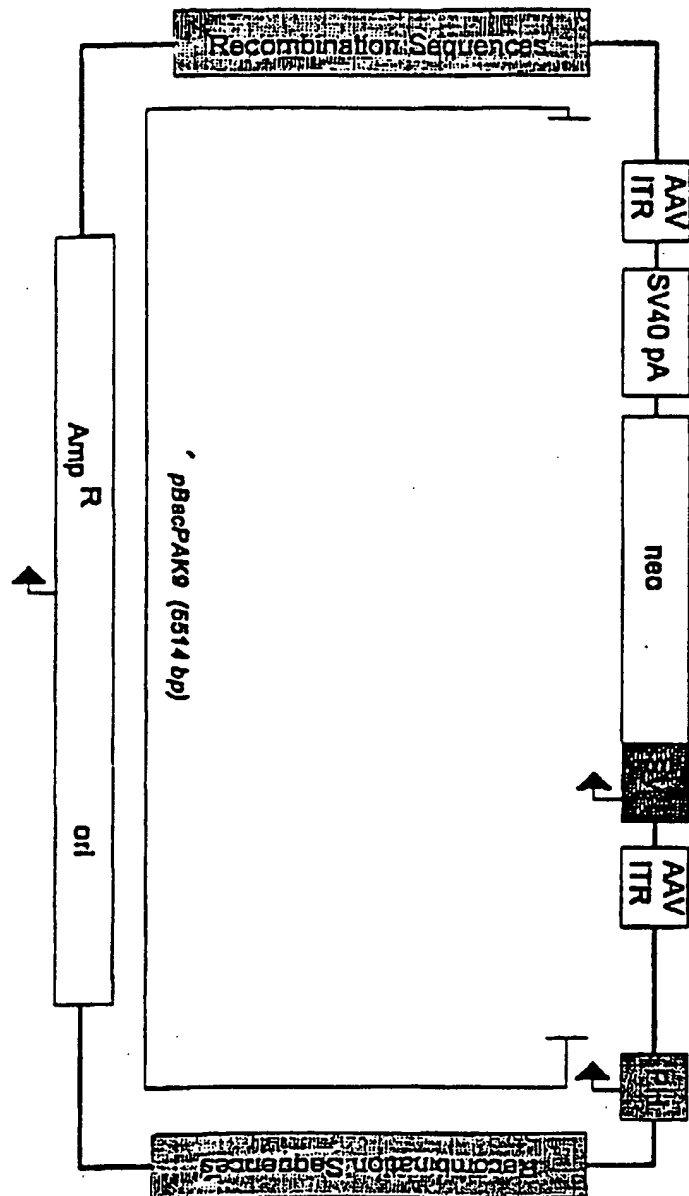
Scheme for auto-excising episome



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FIG. 5

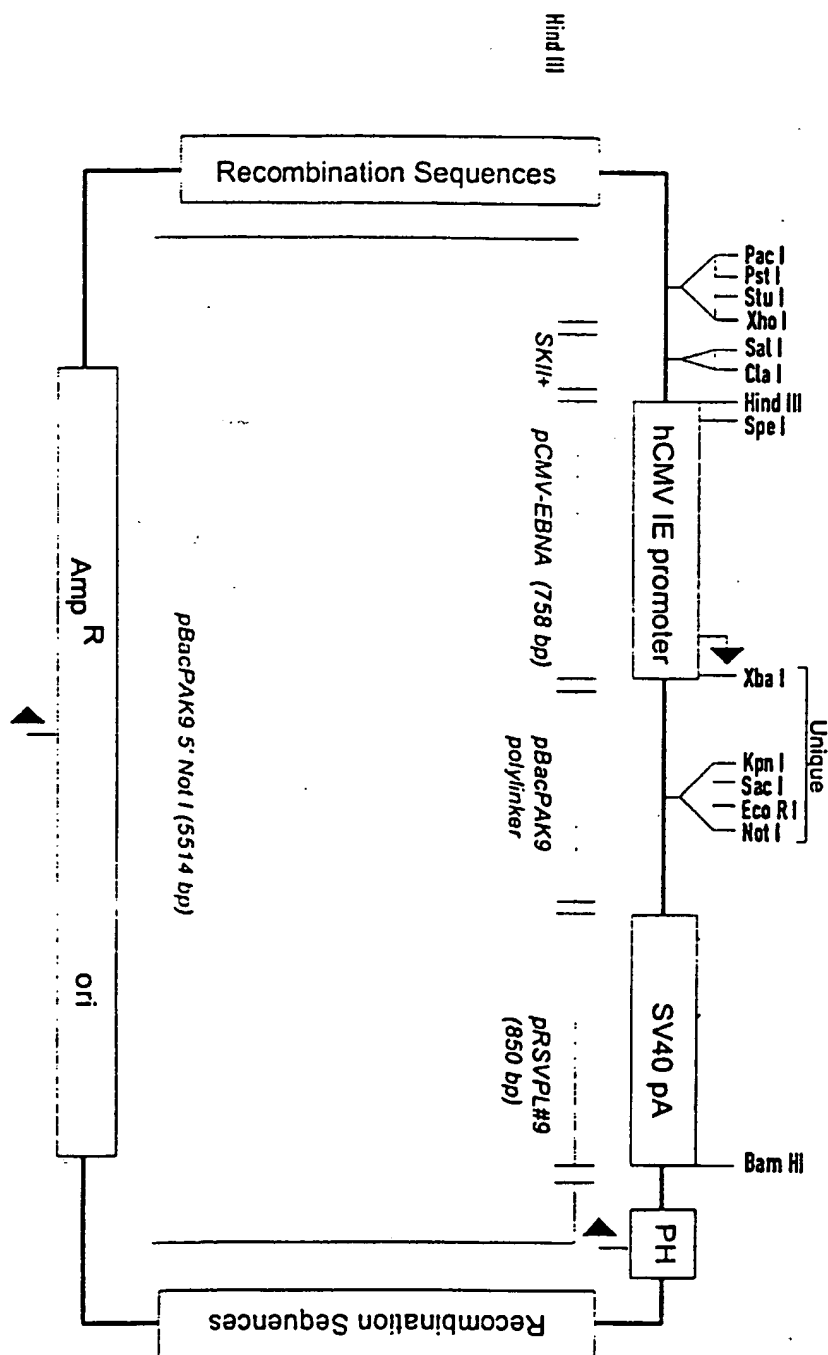
AcMNPV Transfer Plasmid pBV-AVneo



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FIG. 6

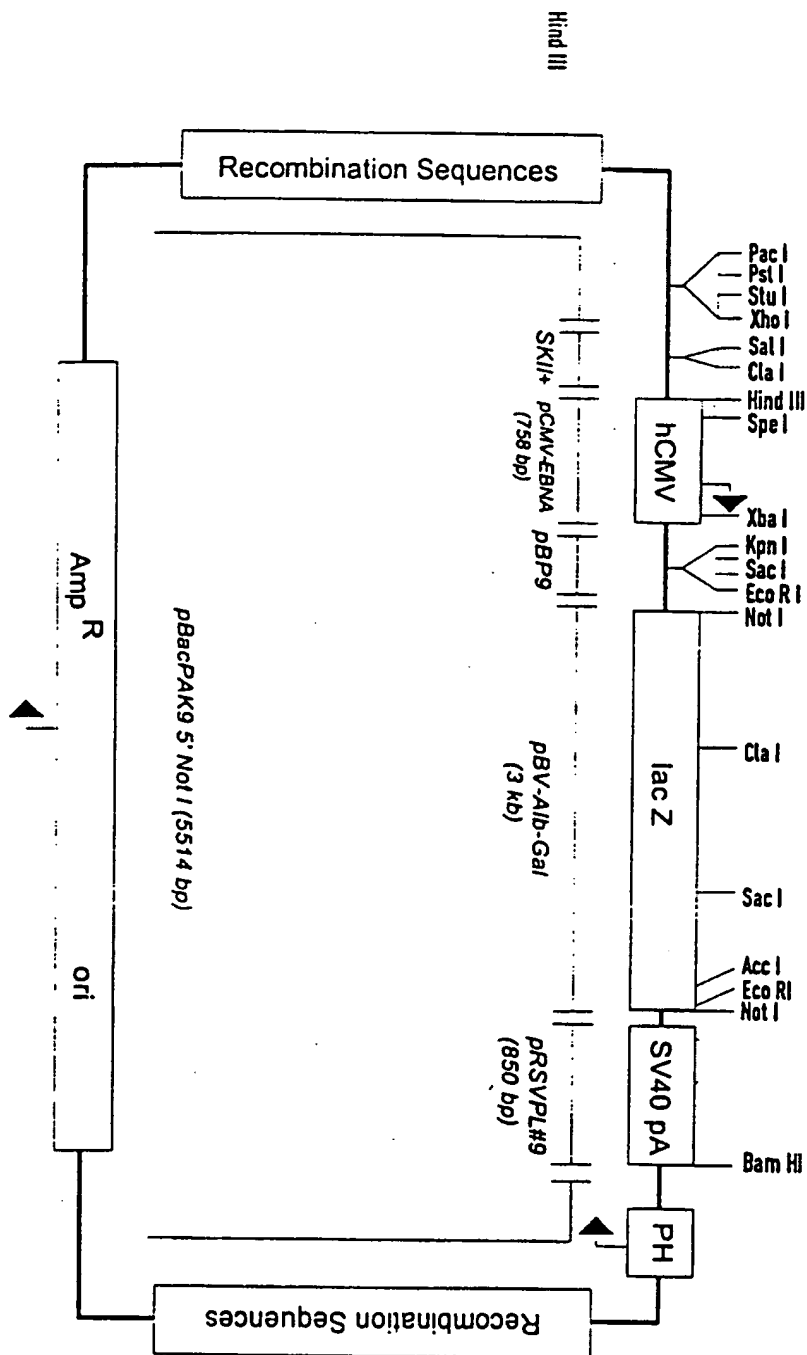
AcMNPV Transfer Plasmid CMV-BV



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FIG. 7

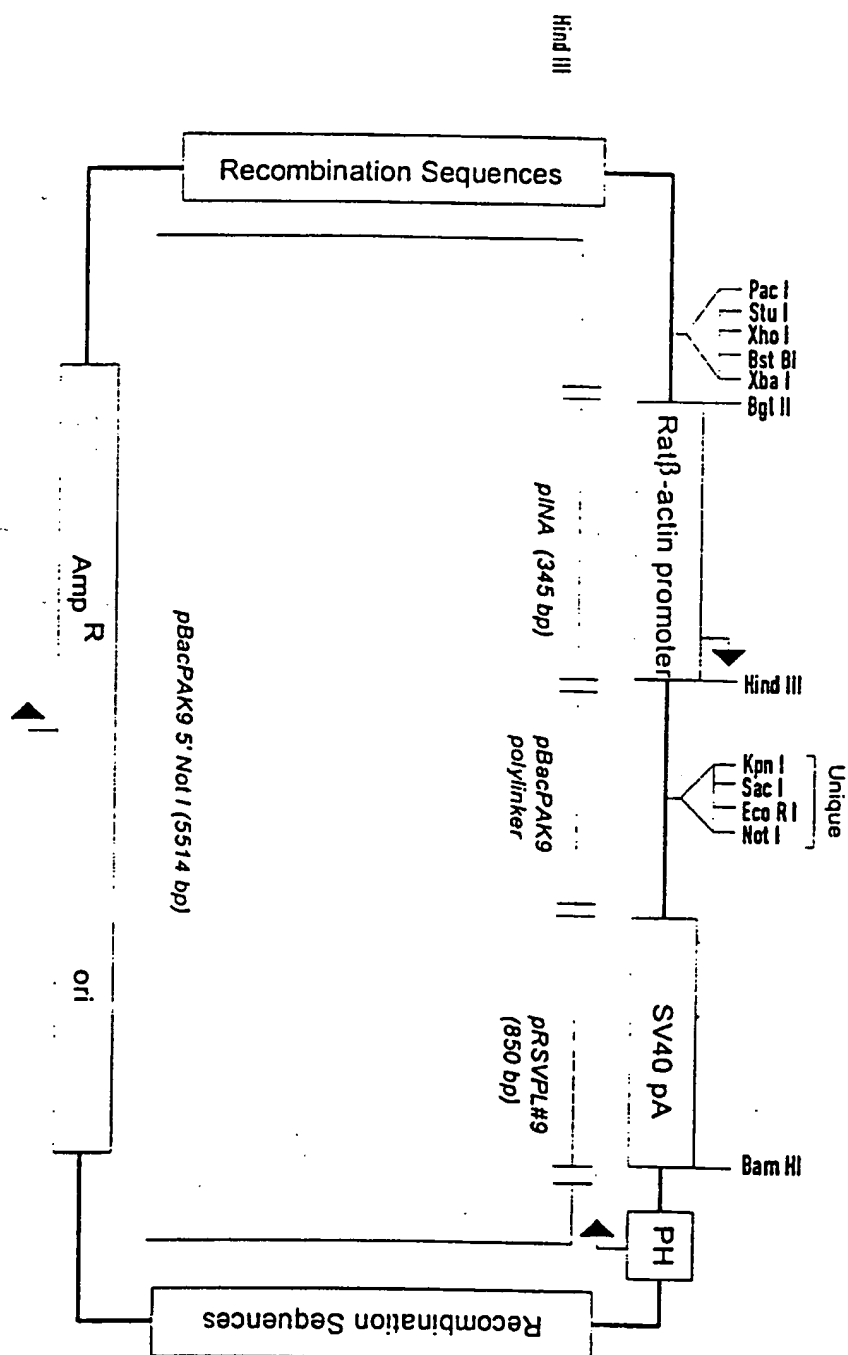
AcMNPV Transfer Plasmid CMVZ-BV



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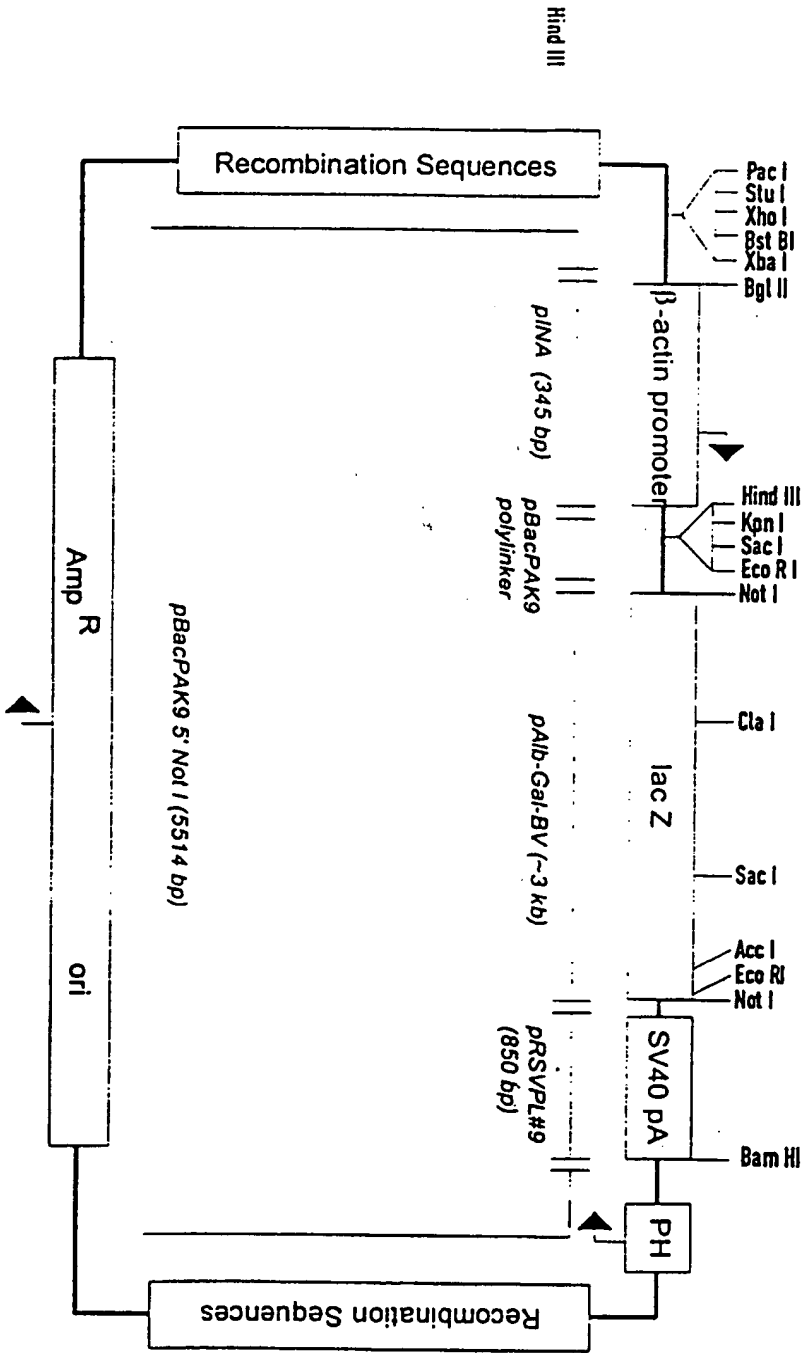
FIG. 8

AcMNPV Transfer Plasmid Act-BV



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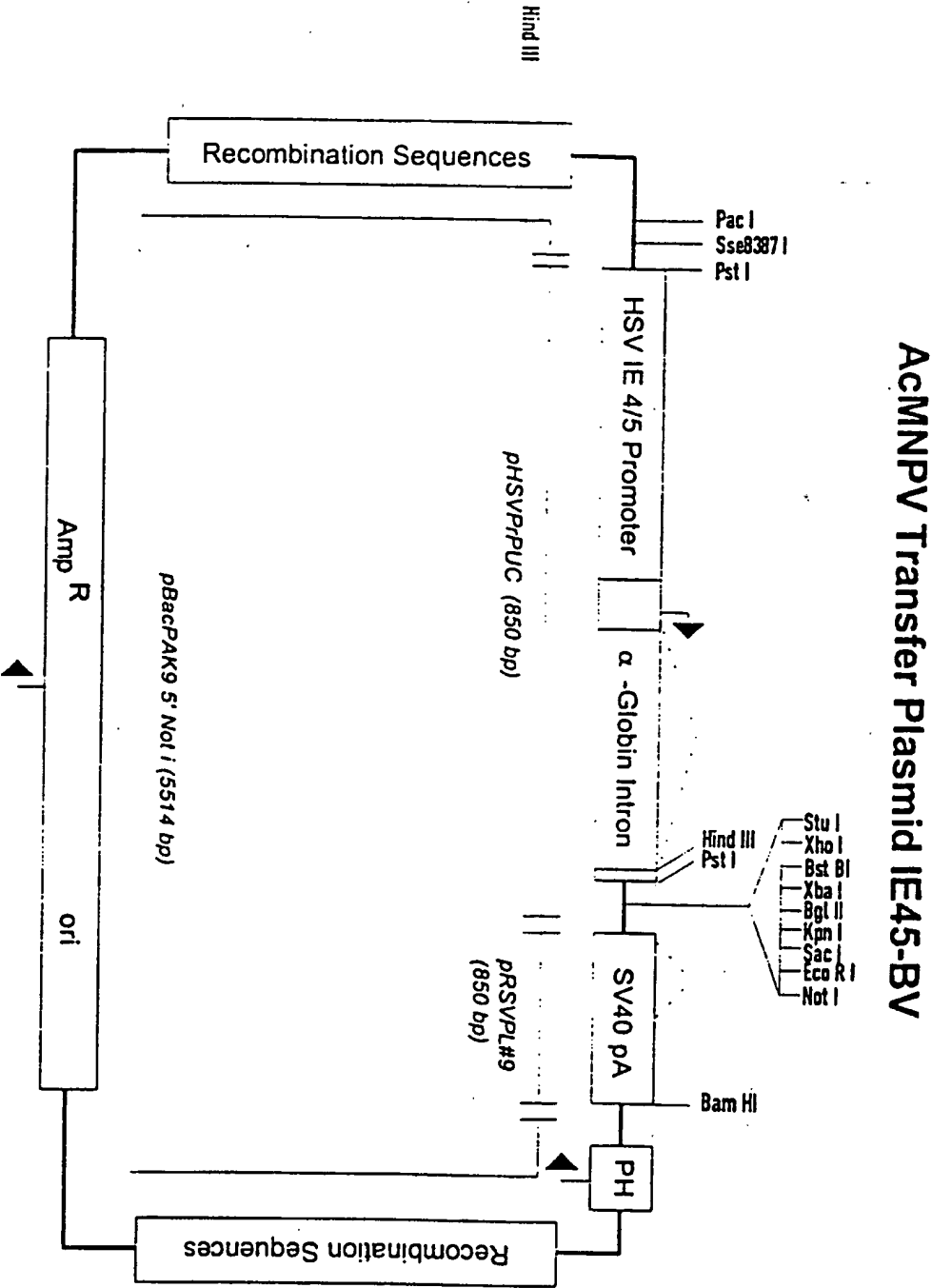
FIG. 9



AcMNPV Transfer Plasmid AZ-BV

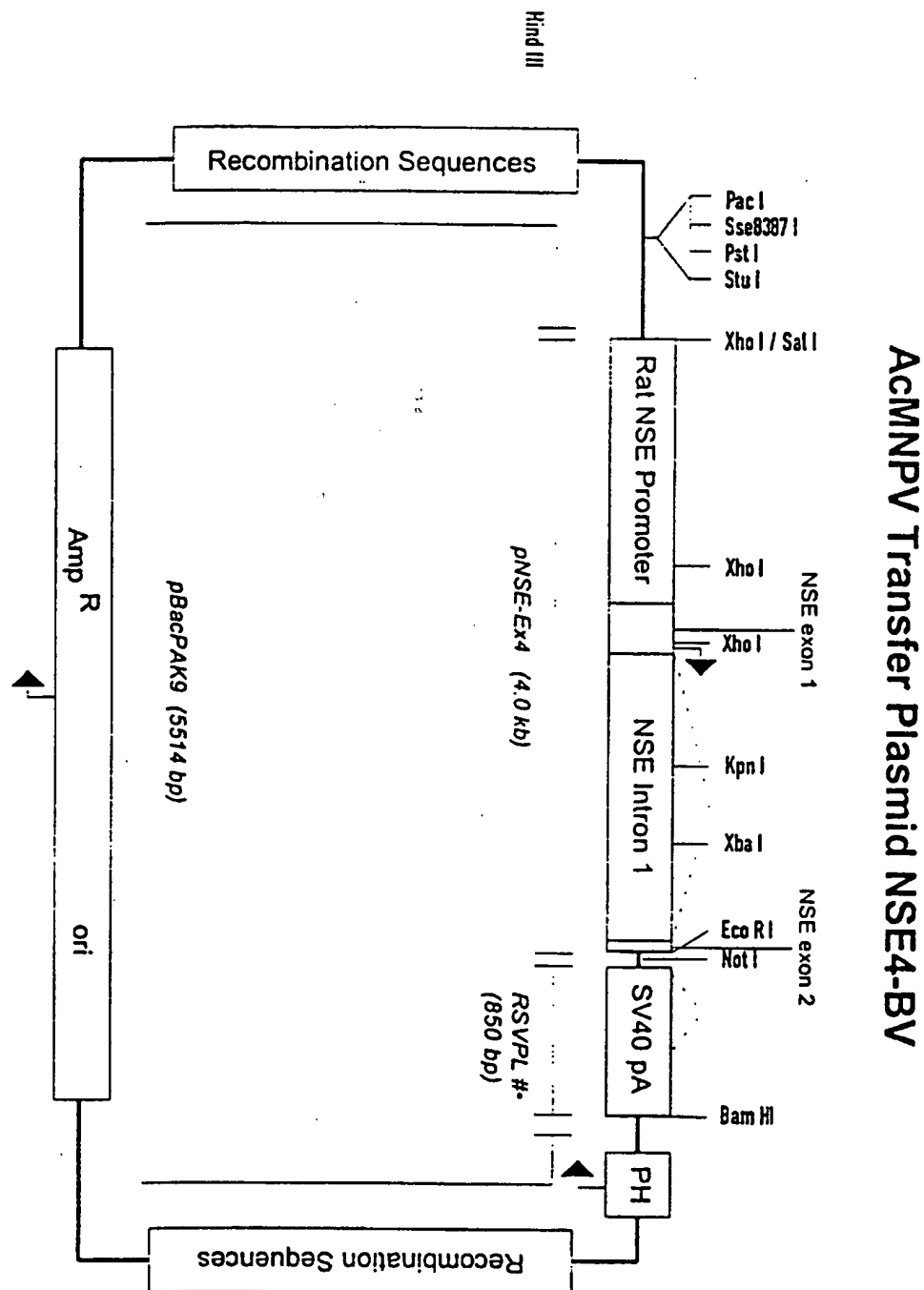
10/31

FIG. 10



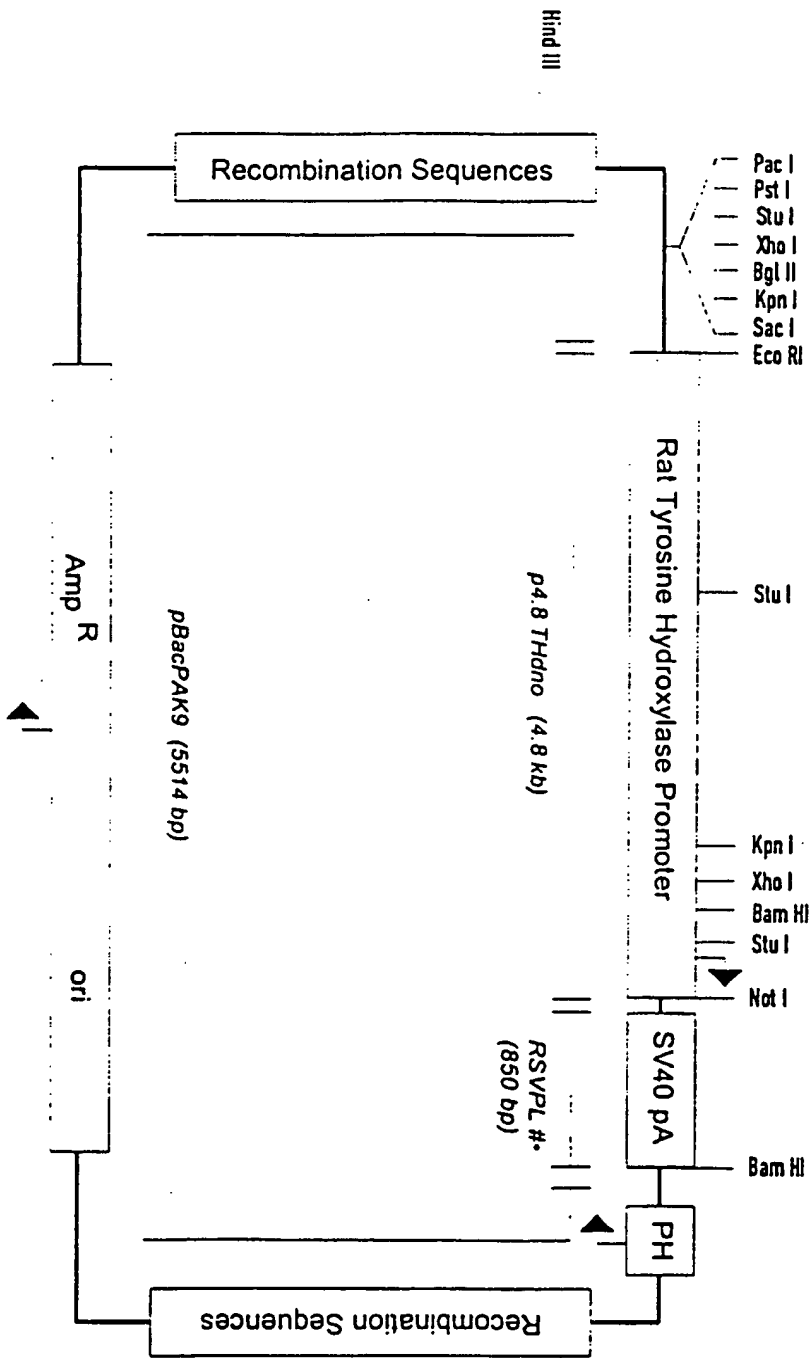
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FIG. 11



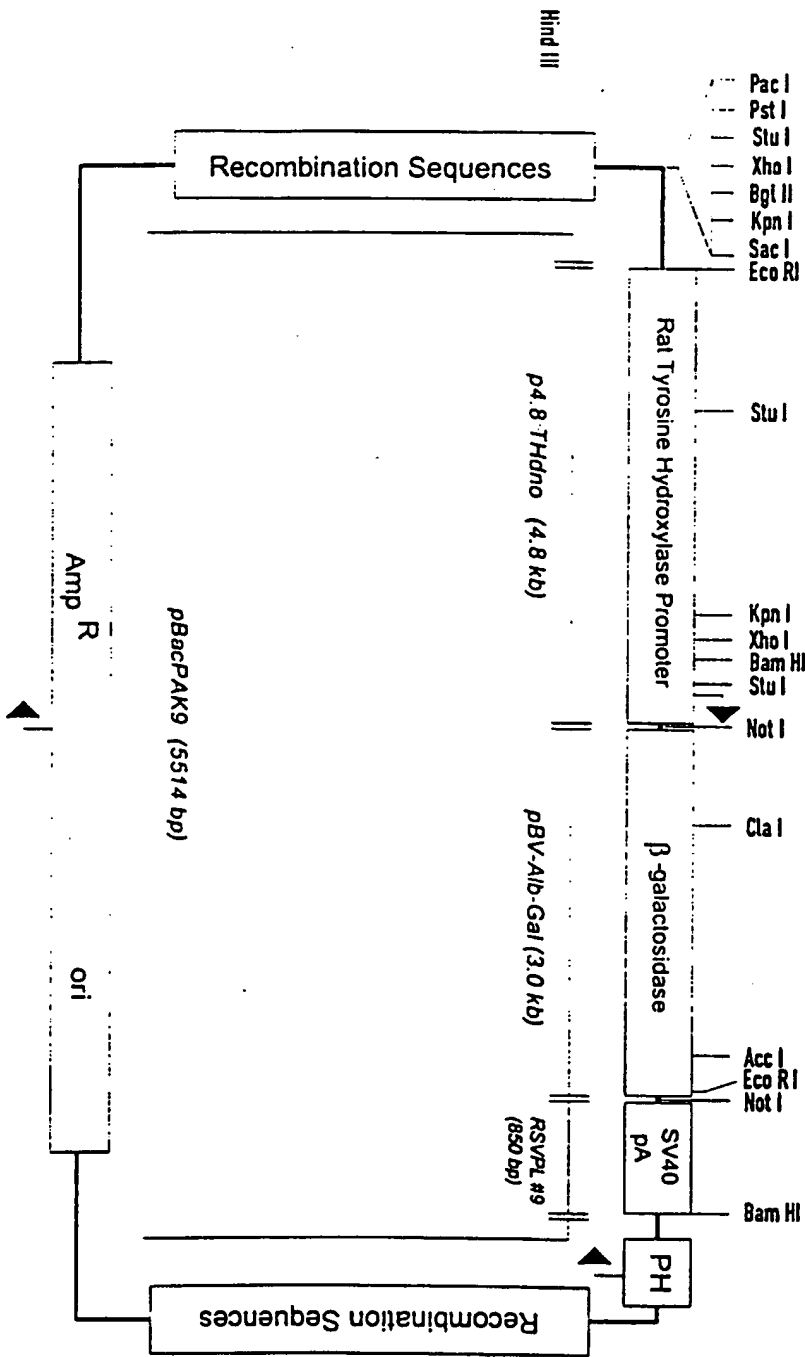
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FIG. 12



AcMNPV Transfer Plasmid TH/SV40/BP9

FIG. 13



AcMNPV Transfer Plasmid TH-Lac/BP9

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FIG. 14A - 14D

14A

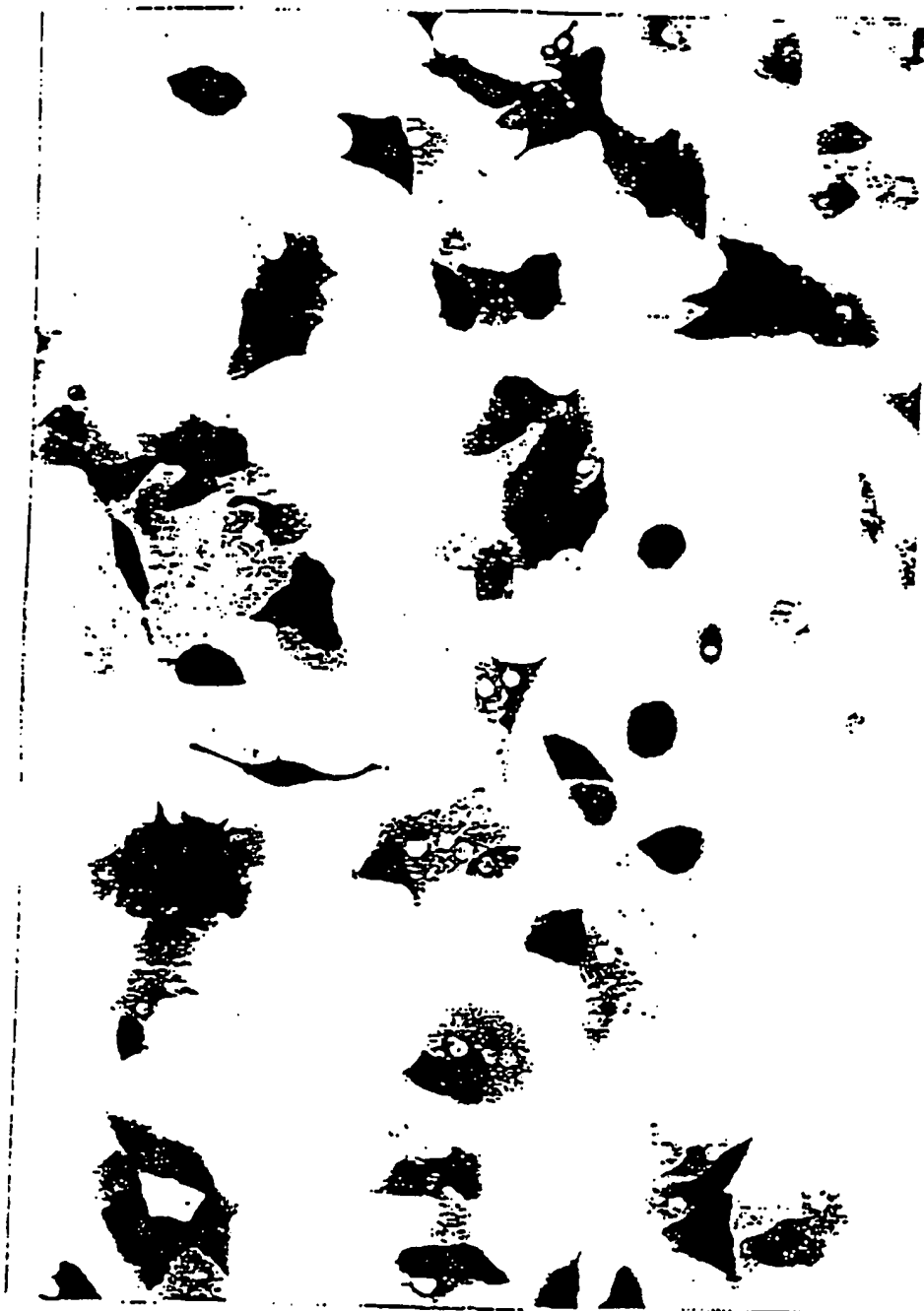
14B

14C

14D

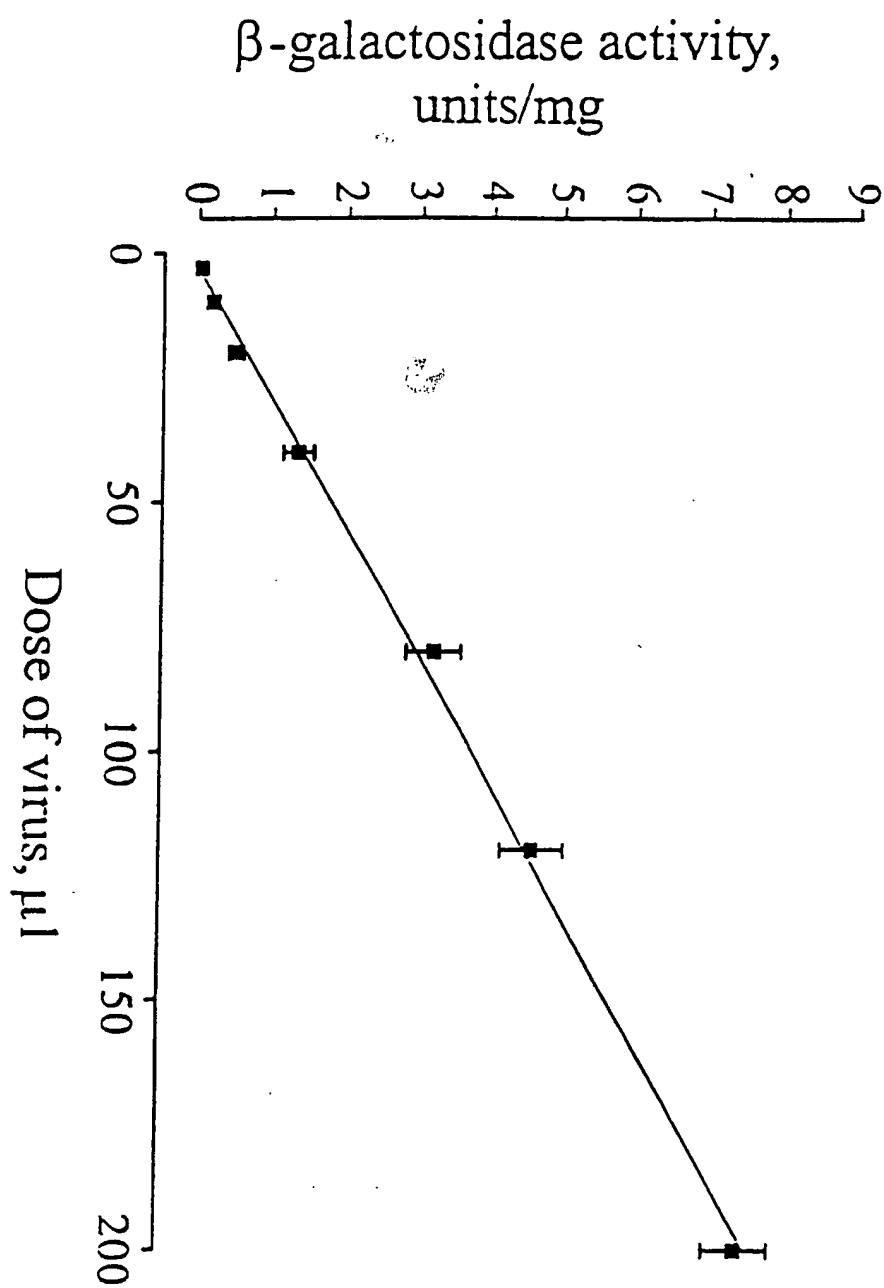
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FIG. 15



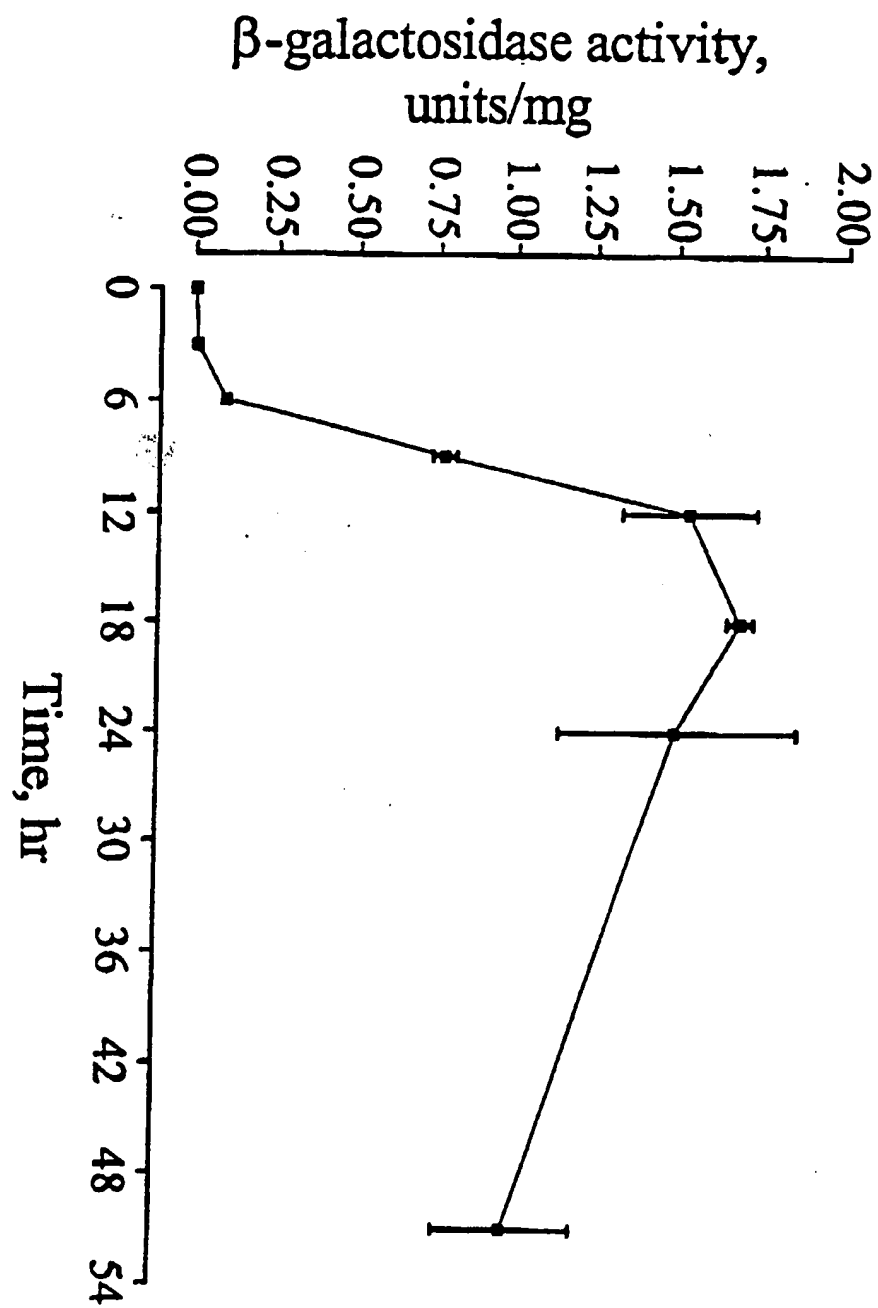
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FIG. 16



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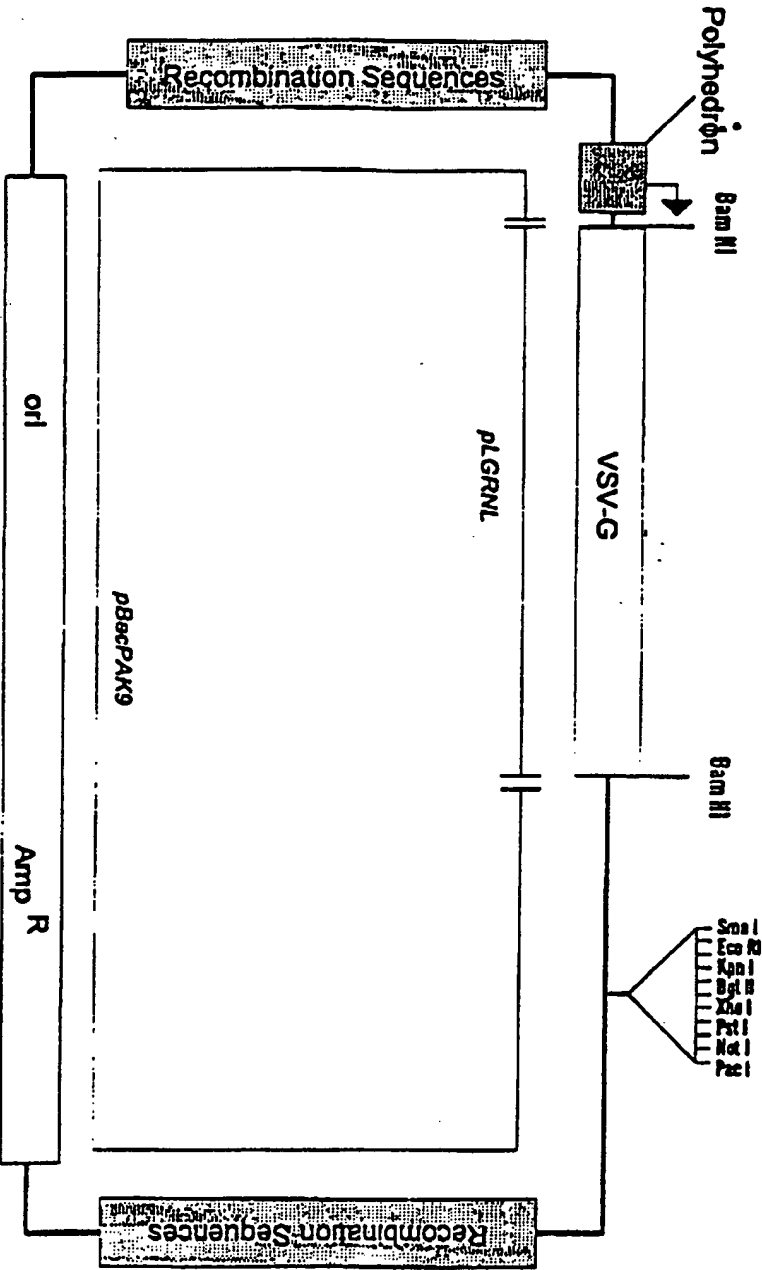
FIG. 17



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FIG. 18

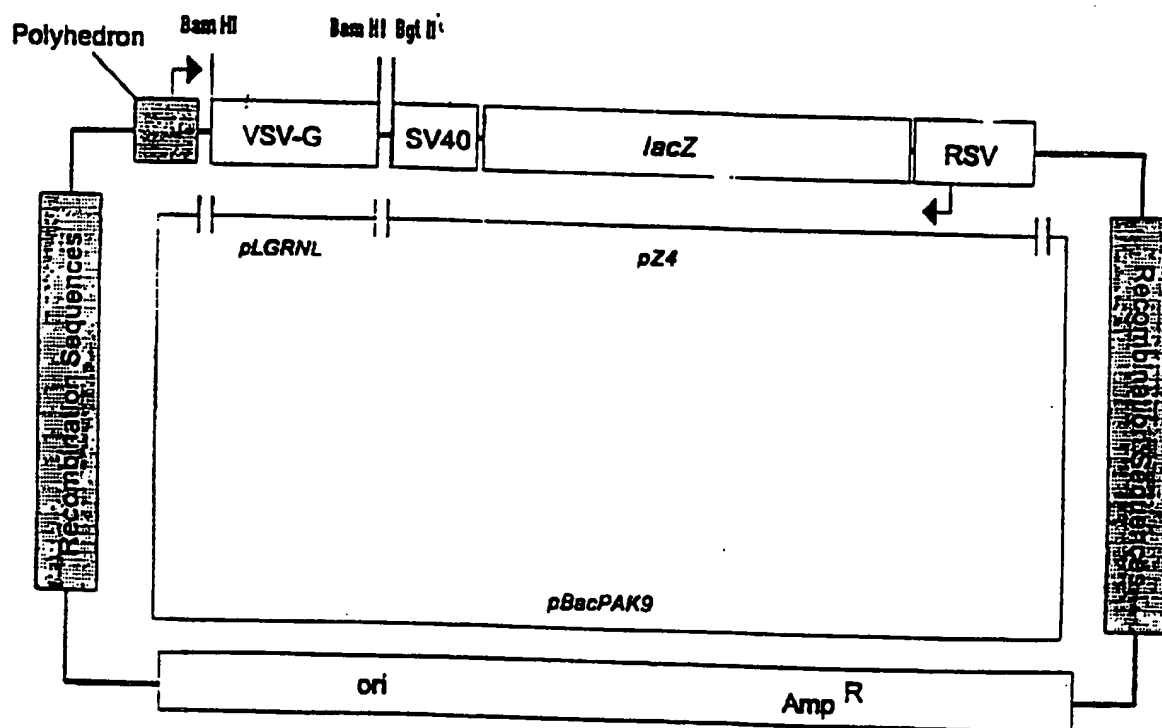
AcMNPV Transfer Plasmid VSV-G/BP9



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FIG. 19

AcMNPV Transfer Plasmid VGZ3

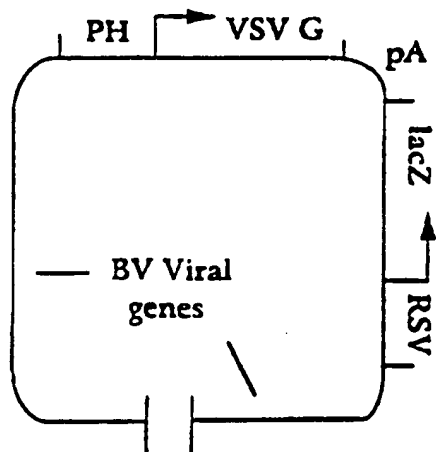


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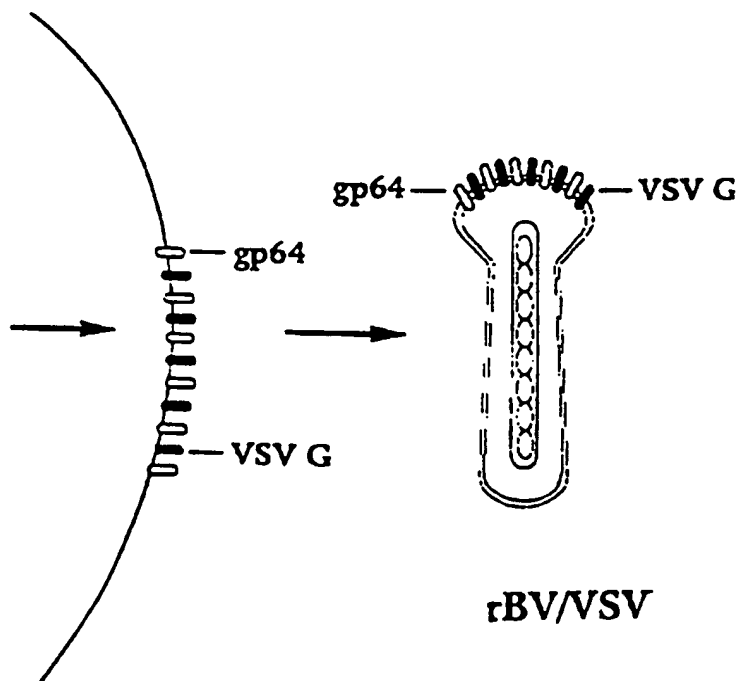
FIG. 20

SECOND GENERATION rBV

VIRUS PACKAGED
IN SF9 CELL



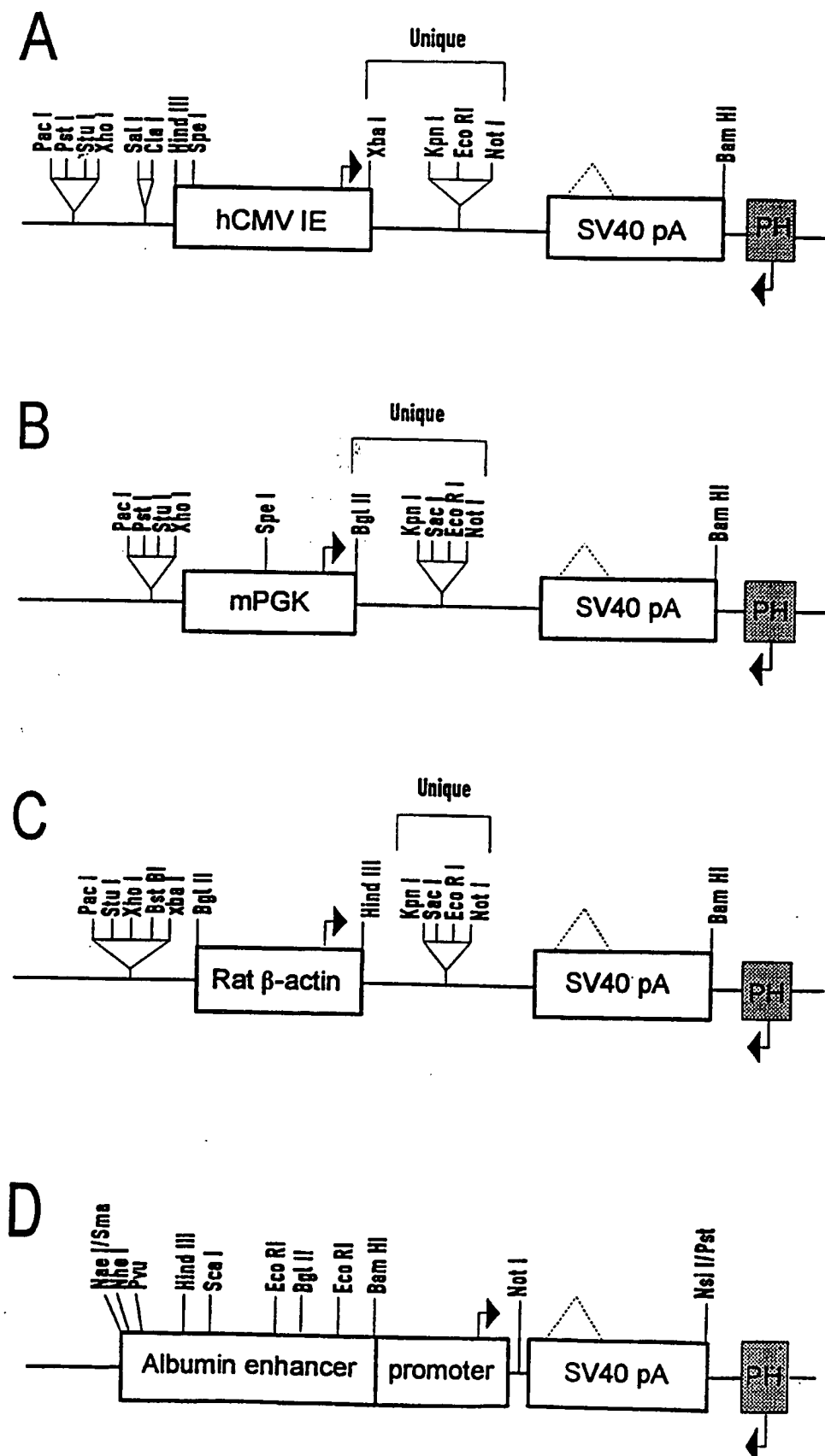
rBV/VSVG GENOME



rBV/VSV

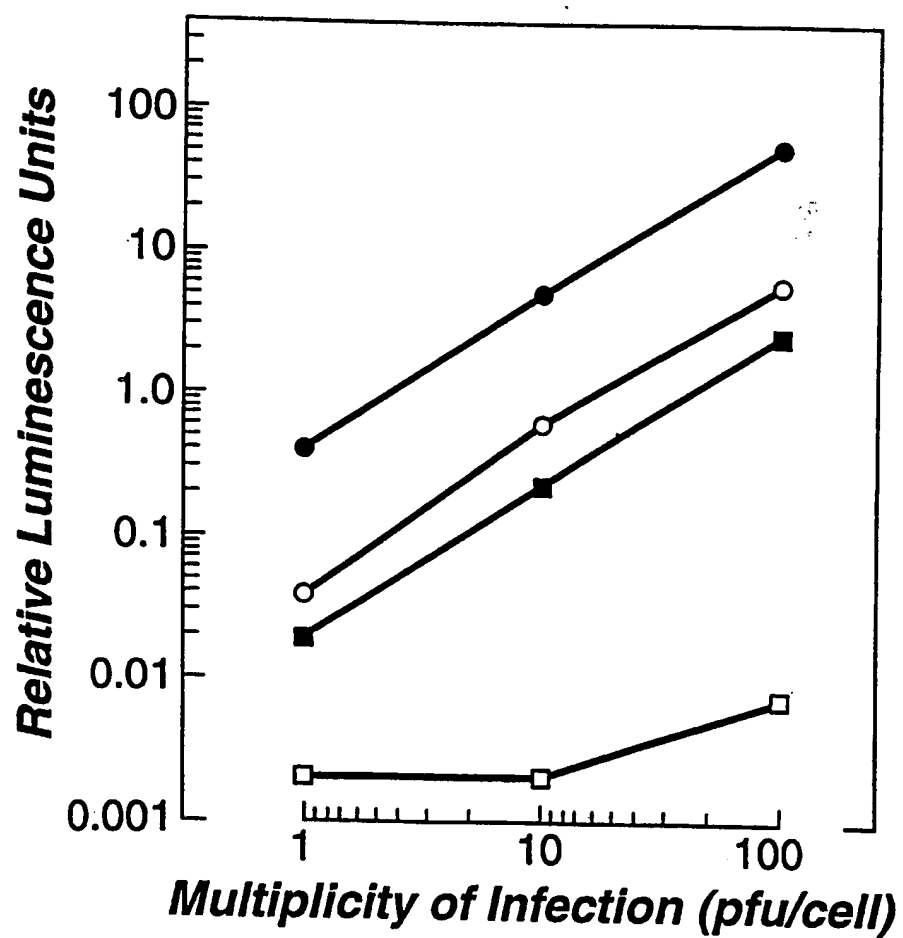
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FIGURE 21



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FIGURE 22



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FIGURE 23

czpgbv.seq Length: 11958

Type: N Check: 7625

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1 AACGGCTCCG CCCACTATTA ATGAAATTAA AAATTCCAAT TTTAAAAAC
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101 GTCGTCGACA TGCTGAACAA CAAGATTAAT ATGCCTCCGT GTATAAAAAA
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1201 AGTATTTTAC TGTTTTCGTA ACAGTTTGT AATAAAAAAA CCTATAAATA
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1301 TTTTATTCA TTGGGTGAA TTGCAAGTTC ACCATAGTTT TTCCACACAA
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5601 CGCCGGTAGC CAGCGCGGAT CATCGGTCAG ACGATTCATT GGCACCATGC

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5651 CGTGGGTTTC AATATTGGCT TCATCCACCA CATACAGGCC GTAGCGGTCG
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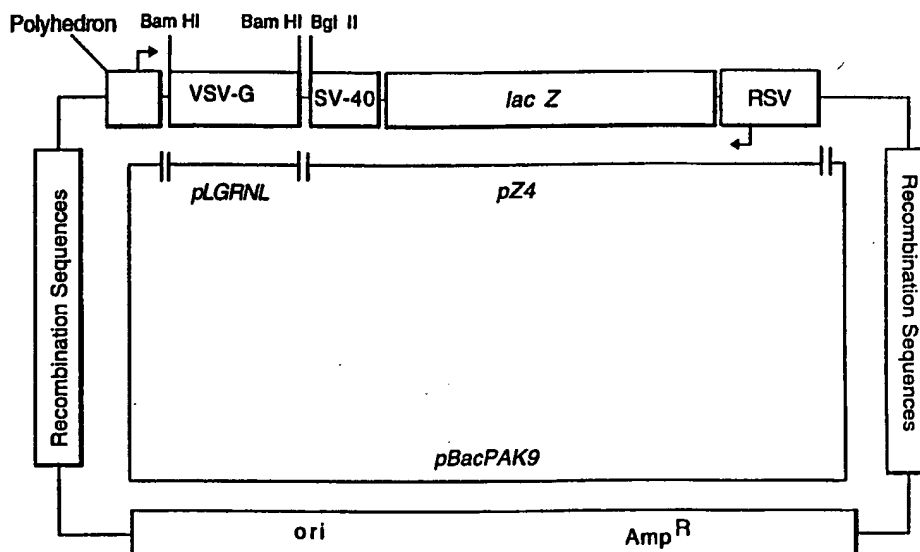
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11951 TAATGCAG



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(54) Title: EXPRESSION OF AN EXOGENOUS GENE IN A MAMMALIAN CELL BY USE OF A NON-MAMMALIAN DNA VIRUS HAVING AN ALTERED COAT PROTEIN



(57) Abstract

Disclosed are methods, nucleic acids, and cells for expressing an exogenous gene in a mammalian cell, involving introducing into the cell a non-mammalian DNA virus (e.g., a baculovirus) having an altered coat protein, the genome of which virus carries an exogenous gene, and growing the cell under conditions such that the gene is expressed. Also disclosed are methods for treating gene deficiency disorders, neurological disorders, or cancers in a mammal by (1) providing to a cell a therapeutically effective amount of a non-mammalian DNA virus having an altered coat protein, the genome of which virus carries an exogenous, therapeutic gene and (2) growing the cell under conditions such that the exogenous gene is expressed in the mammal.

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BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	NZ	New Zealand		
CM	Cameroon			PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakhstan	RO	Romania		
CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	SG	Singapore		

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 97/16041

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C12N15/86 C12N15/62 C12N15/47 C12N7/01 C12N5/10
 C07K14/145 A61K48/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C12N C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 96 09074 A (GEN HOSPITAL CORP) 28 March 1996 see abstract see page 9, line 15 - line 16 see page 18, line 35 - page 19, line 5 see page 54, line 10 - line 12 see claims 1-7, 26-34, 39-43, 51-56, 59 ---	1-52
Y	WO 92 14829 A (UNIV CALIFORNIA ;VIAGENE INC (US)) 3 September 1992 see abstract see page 5 -----	1-52

☐ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

* Special categories of cited documents :

A document defining the general state of the art which is not considered to be of particular relevance

E earlier document but published on or after the international filing date

L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

O document referring to an oral disclosure, use, exhibition or other means

P document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

Z document member of the same patent family

Date of the actual completion of the international search

9 March 1998

Date of mailing of the international search report

17. 03. 98

Name and mailing address of the ISA

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Authorized officer

Galli, I

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 97/16041

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
see FURTHER INFORMATION sheet PCT/ISA/210
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International Application No. PCT/US 97/16041

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Remark : Although claims 5-20 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 97/16041

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9609074 A	28-03-96	AU 3675095 A	09-04-96
		CA 2200835 A	28-03-96
		EP 0785803 A	30-07-97
		ZA 9507797 A	08-07-96

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		US 5670354 A	23-09-97
